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(54) **Vitamin D3 analogs**

Vitamin D3 Analoge

Analogues de la vitamine D3

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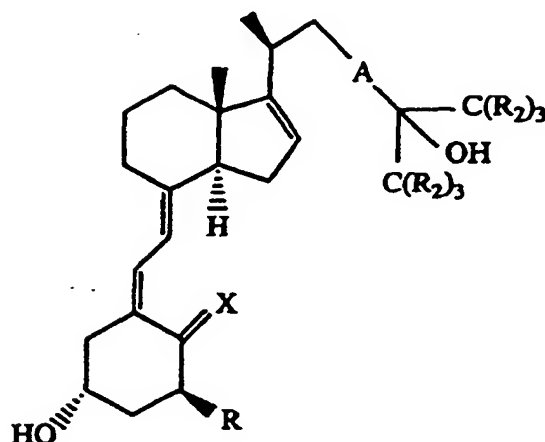
• E. ELSTNER ET AL: CANCER RESEARCH, vol.
55, 1 July 1995, pages 2822-2830, XP002078809

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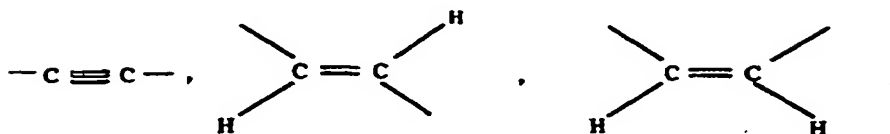
EP 0 808 833 B1

Description

[0001] The invention relates to Vitamin D₃ analogs, particularly 20epi-16-ene analogs of Vitamin D₃ of the formula



wherein R is fluorine, R₂ is hydrogen, lower alkyl or halogen, X is =CH₂, and A is



or

-CH₂-CH₂-, provided that when A is -CH₂-CH₂-, R₂ is hydrogen or lower alkyl.

[0002] As used herein, "lower alkyl" denotes a straight or branched chain alkyl group containing 1 to 4 carbon atoms, for example, methyl, ethyl, propyl, isopropyl, butyl or t-butyl. Preferably lower alkyl is methyl or ethyl. Halogen means fluorine, iodine, bromine or chlorine, preferably, fluorine.

[0003] The compounds of formula I differ from known compounds by the presence of a fluorine atom in position 1 instead of hydroxy. Examples of such 1-hydroxy compounds are known from EP 325279 (having "natural" configuration in position 20); Cancer Research 1995, 55, 2822-2830 (having "epi" configuration in position 20); and from WO 92/03414 and JP 7188281 (having saturation in position 16).

[0004] Compounds of formula I induce differentiation and inhibition of proliferation in various skin and cancer cell lines. Accordingly, the compounds of formula I are useful as agents for the treatment of hyperproliferative skin diseases, such as, psoriasis. Compounds of formula I are also useful in the treatment of neoplastic diseases, such as, leukemia or breast cancer and sebaceous gland diseases, such as, acne or seborrheic dermatitis. Compounds of formula I, particularly 1α-fluoro-25-hydroxy-16,23E-diene-26,27-bishomo-20-epi-cholecalciferol, are useful in the treatment of osteoporosis.

[0005] Furthermore, when applied topically to the skin of a patient the compounds of formula I reverse the conditions associated with photodamage. Hence, by the topical application of compounds of formula I to the skin of patients which has been damaged through sun exposure, the effects of wrinkling, elastosis and premature aging can be reversed leading to an improvement in the appearance of the skin. In other words, through the topical administration of the compounds of the formula I, the acceleration of repair of dermal damage is accomplished so as to provide the skin with a smoother and younger appearance.

[0006] The invention also relates to a pharmaceutical composition comprising a compound of formula I and a pharmaceutically acceptable carrier. Such compositions can be used in inducing differentiation and inhibition of proliferation in skin and cancer cell lines, particularly for treating hyperproliferative skin diseases, such as psoriasis; neoplastic diseases, such as leukemia; and sebaceous gland diseases, such as acne; as well as in reversing the conditions associated with photodamage, particularly for the topical treatment of the skin damaged through sun exposure, the

effects of wrinkling, elastosis and premature aging. The invention also relates to the use of the compounds of formula I in the manufacture of medicaments for treating the above mentioned diseases and conditions. The invention also relates to a process for preparing compounds of formula I and intermediates of formula XII.

[0007] In a compound of formula I, R₂ is preferably hydrogen or fluorine. Preferred compounds of formula I are:

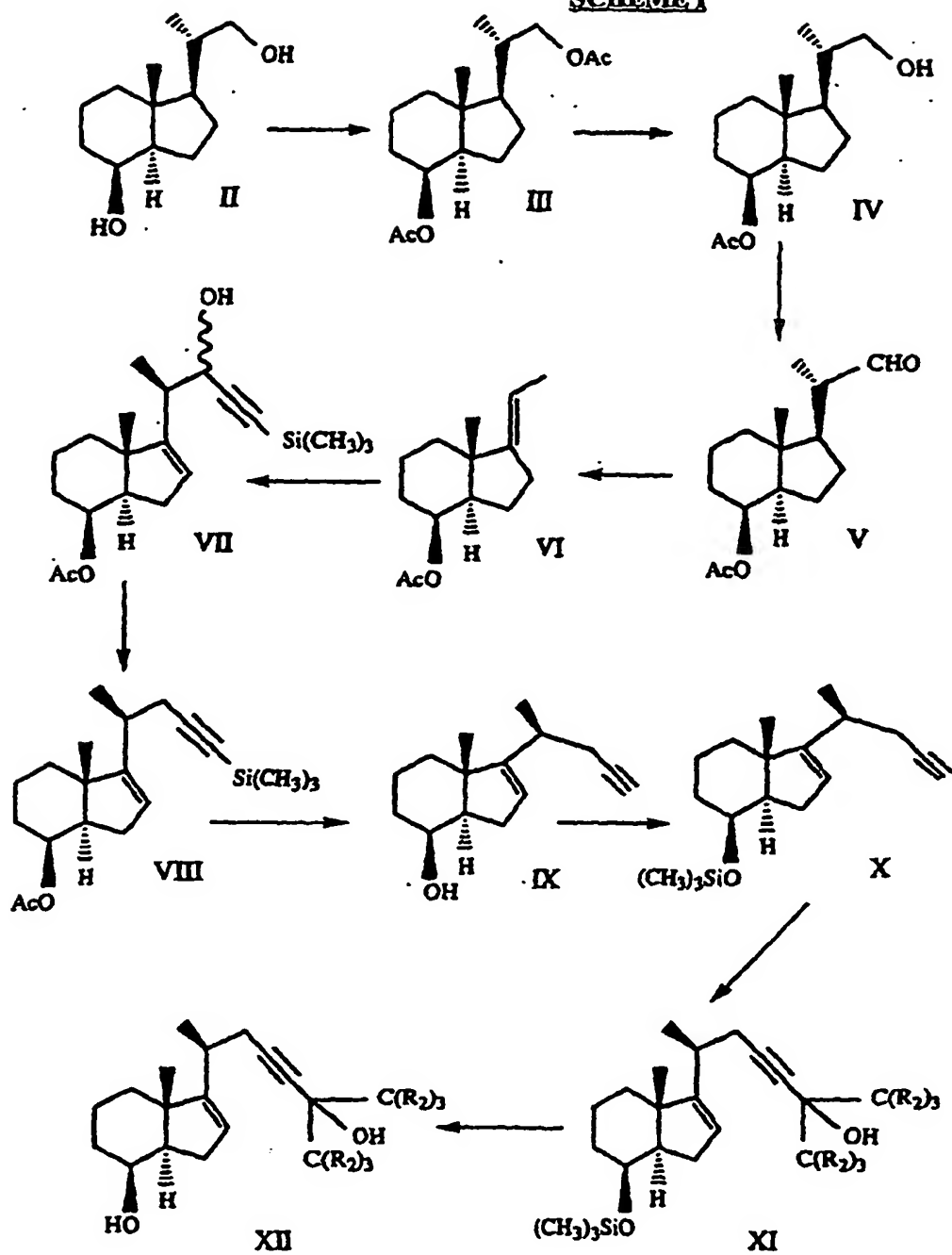
1 α -fluoro-25-hydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-cholecalciferol;

1 α -fluoro-25-hydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol.

[0008] The compounds of formula I are prepared as hereafter described in Schemes I-V and the Examples.

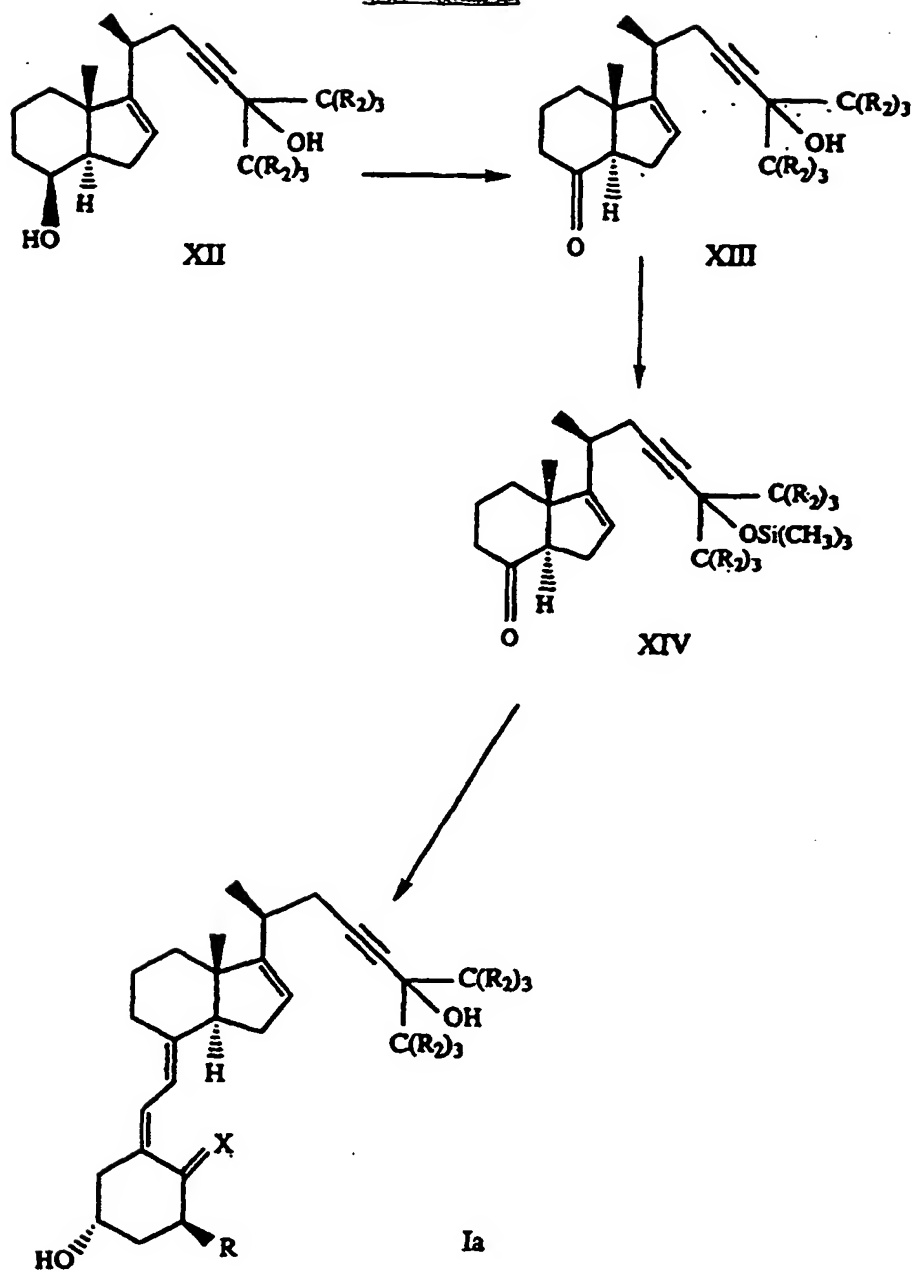
[0009] The last step in the process for preparing the compounds of formula I comprises deprotecting corresponding compounds containing silyl protected hydroxy groups.

SCHEME I



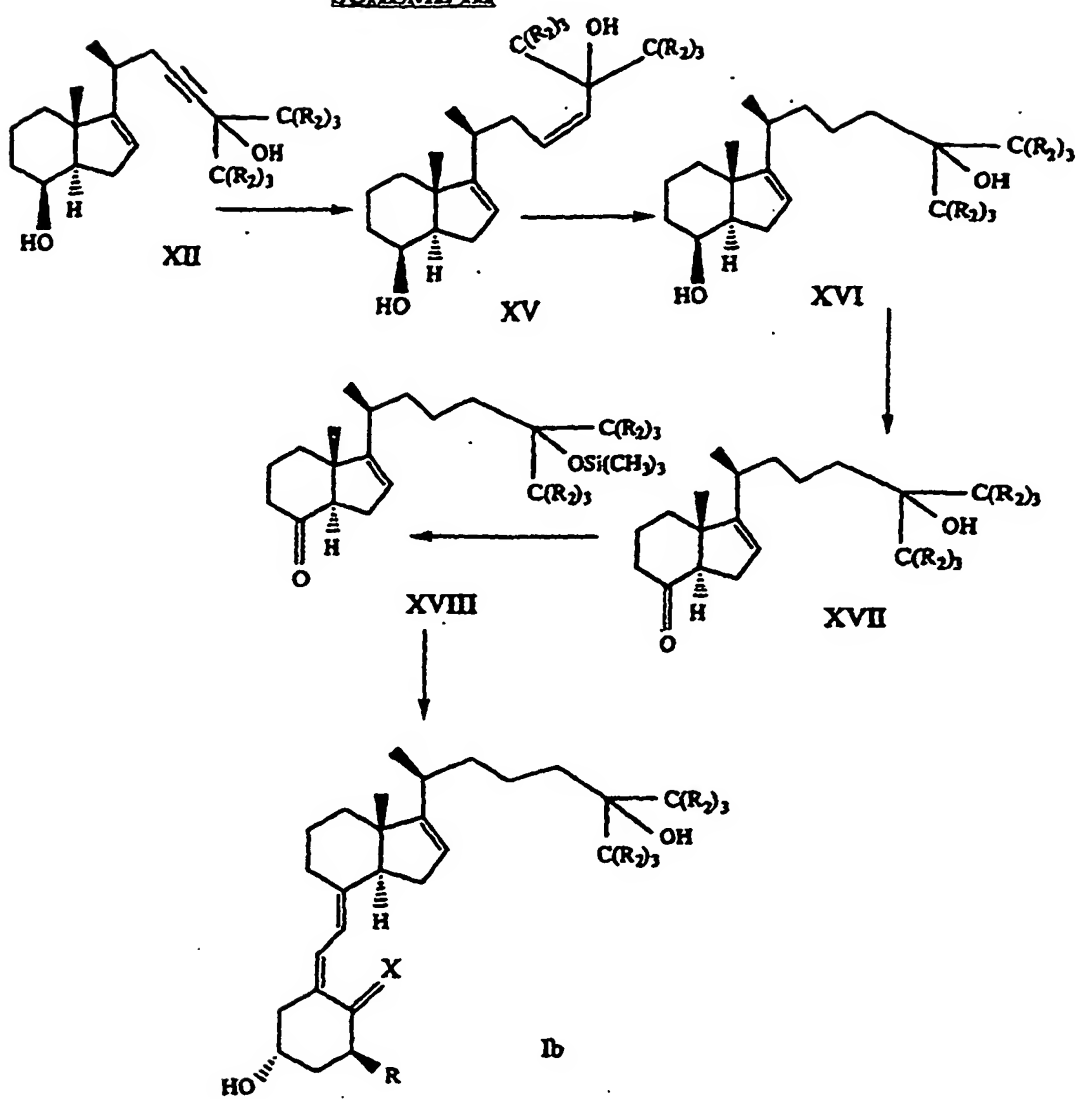
wherein R_2 is as described above.

SCHEME II

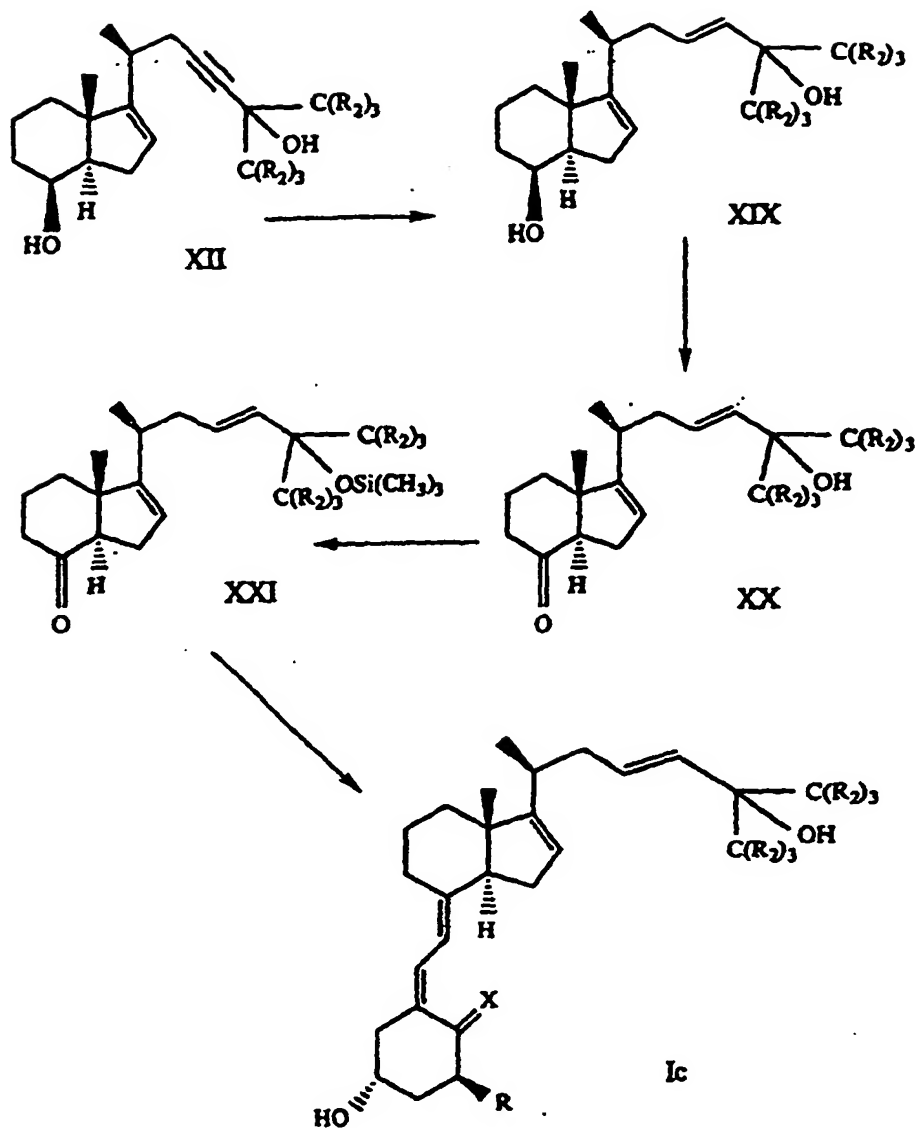


wherein R, X and R_2 are as described above.

SCHEME III

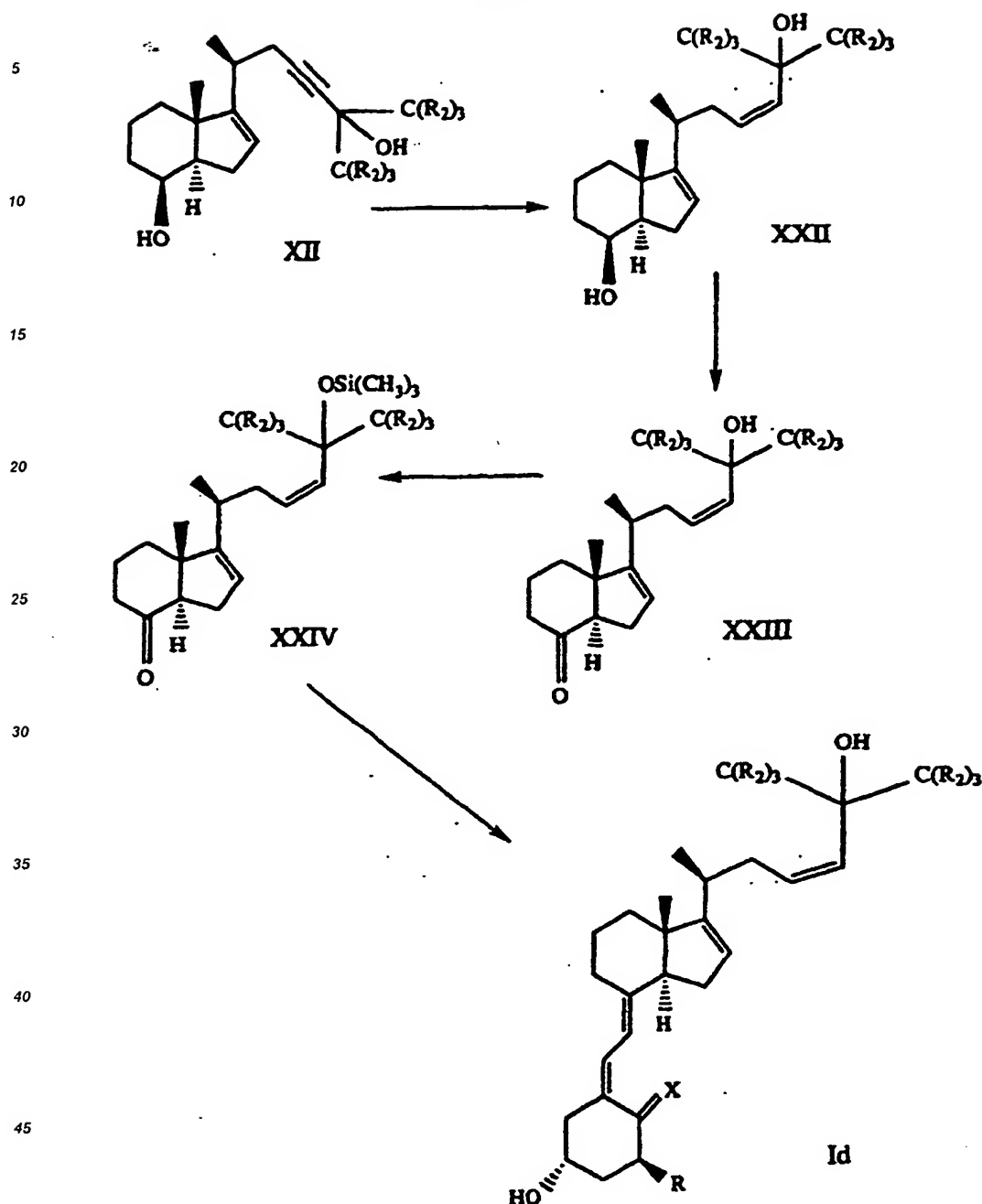


wherein R, X are as described above and R_2 is hydrogen or lower alkyl.

SCHEME IV

wherein R, X and R_2 are as described above.

SCHEME V



wherein R, X and R₂ are as described above.

[0010] In above Scheme I, the compound of formula II, a known compound (B. M. Trost, P.R. Bernstein, P.C. Funf-schilling, J. American Chemical Society 101, 4378 (1979)), is converted to the compound of formula III by reaction with acetic anhydride, pyridine and dimethylaminopyridine in a chlorinated hydrocarbon solvent, such as dichloromethane. The reaction is carried out at 0 C to 50 C, preferably room temperature, preferably under an argon atmosphere.

[0011] The compound of formula III is converted to the compound of formula IV by reaction with a base, such as, sodium carbonate in an alcohol solvent, such as methanol, under, preferably, an argon atmosphere.

[0012] The compound of formula IV is converted to the compound of formula V by reaction with oxalyl chloride and dimethylsulfoxide in a chlorinated hydrocarbon solvent, such as, dichloromethane, under an argon atmosphere.

[0013] The compound of formula V is converted to the compound of formula VI by reaction with benzalacetone in the presence of palladium on charcoal catalyst.

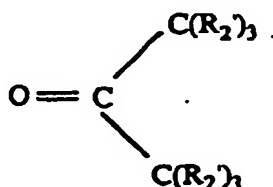
[0014] The compound of formula VI is converted to the compound of formula VII by reaction with a 3-trimethylsilyl-propynal and a Lewis acid such as dimethylaluminum chloride in a chlorinated hydrocarbon solvent, such as dichloromethane.

[0015] The compound of formula VII is converted to the compound of formula VIII by reaction of the corresponding phenylthiono-carbonate with tri-n-butyltin hydride and triethylborane in toluene.

[0016] The compound of formula VIII is converted to the compound of formula IX by reaction with a base, such as, sodium hydroxide in an alcohol solvent, such as ethanol. The reaction is carried out at a temperature in the range of 50 C to 100 C, preferably 80 C.

[0017] The compound of formula IX is converted to the compound of formula X by reaction with 1-(trimethylsilyl)imidazole in a chlorinated hydrocarbon solvent, such as anhydrous methylene chloride.

[0018] The compound of formula X is converted to a compound of formula XI by reaction with the corresponding compound of formula



wherein R_2' is hydrogen, or lower alkyl in the presence of a base such as n-butyllithium.

The reaction is preferably carried out at -78 C.

[0019] When the compounds of formula I wherein R_2 is halogen are prepared, the compound of formula X is reacted with a halogenated acetone, such as hexafluoroacetone in the presence of a base such as n-butyllithium.

[0020] The compound of formula XI is converted to the corresponding compound of formula XII by reaction with tetrabutylammonium fluoride in an ether solvent such as, tetrahydrofuran.

[0021] As set forth in Scheme II, a compound of formula XII is converted to a corresponding compound of formula XIII by reaction with an oxidizing agent such as, pyridinium dichromate in a chlorinated hydrocarbon solvent such as, anhydrous methylene chloride.

[0022] A compound of formula XIII is converted to a corresponding compound of formula XIV by reaction with 1-(trimethylsilyl)imidazole in a chlorinated hydrocarbon solvent, such as, anhydrous methylene chloride.

[0023] A compound of formula XIV is converted to the corresponding compound of formula Ia wherein R is fluorine by reaction with {3S-(3 α ,5 β ,Z)}-2-[2-[2-methylene-3-fluoro-5-[[1,1-dimethylethyl]dimethylsilyl]oxy]cyclohexylidene]ethyl]diphenylphosphine oxide in a ether like solvent such as tetrahydrofuran in the presence of n-butyllithium as a base.

[0024] In the above Scheme III, a compound of formula XII is reduced to a corresponding compound of formula XV by reaction with hydrogen and Lindlar catalyst in an organic solvent, such as, a combination of ethyl acetate, hexane and ethanol, in the presence of quinoline.

[0025] A compound of formula XV is converted to a corresponding compound of formula XVI by reaction with hydrogen in the presence of a catalyst such as 1,4-bis(diphenylphosphino)butane 1,5-cyclooctadiene rhodium tetrafluoroborate and mercury in a chlorinated hydrocarbon solvent, such as, methylene chloride.

[0026] A compound of formula XVI is oxidized to a corresponding compound of formula XVII by reaction with pyridinium dichromate in a chlorinated hydrocarbon solvent, such as methylene chloride.

[0027] A compound of formula XVII is converted to a corresponding compound of formula XVIII by reaction with 1-(trimethylsilyl)imidazole in a chlorinated hydrocarbon solvent, such as methylene chloride.

[0028] A compound of formula XVIII is converted to the corresponding compound of formula Ib wherein R is fluorine by reaction with {3S-(3 α ,5 β ,Z)}-2-[2-[2-[methylene-3-fluoro-5-[[1,1 dimethylethyl]dimethylsilyl]oxy]cyclohexylidene]ethyl]diphenylphosphine oxide in the presence of n-butyllithium as a base in anhydrous tetrahydrofuran as a solvent, at a temperature of -78 C.

[0029] In above Scheme IV, a compound of formula XII is converted to a corresponding compound of formula XIX by reaction with a reducing agent such as, lithium aluminum hydride in an ether solvent, such as, tetrahydrofuran in the presence of sodium methoxide as a base.

[0030] A compound of formula XIX is converted to a corresponding compound of formula XX by reaction with an oxidizing agent such as, pyridinium dichromate in a chlorinated hydrocarbon solvent, such as, methylene chloride.

[0031] A compound of formula XX is converted to a corresponding compound of formula XXI by reaction with 1-(trimethylsilyl)imidazole in a chlorinated hydrocarbon solvent, such as, methylene chloride.

[0032] A compound of formula XXI is converted to a corresponding compound of formula Ic wherein R is fluorine by reaction with [3S-(3 α ,5 β ,Z)]-2-[2-methylene-3-fluoro-5-[[[(1,1-dimethylethyl)dimethylsilyl]oxy]cyclohexylidene]ethyl]diphenylphosphine oxide in a ether like solvent, such as tetrahydrofuran, in the presence of a base such as n-butyllithium.

[0033] As set forth in Scheme V above, a compound of formula XII is converted to a corresponding compound of formula XXII by hydrogenation with Lindlar catalyst in the presence of quinoline in a mixture of solvents, such as a combination of ethyl acetate, hexane and ethanol.

[0034] A compound of formula XXII is converted to a corresponding compound of formula XXIII by reaction with oxidizing agent, such as, pyridinium dichromate in a chlorinated hydrocarbon solvent, such as, methylene chloride.

[0035] A compound of formula XXIII is converted to a corresponding compound of formula XXIV by reaction with 1-(trimethylsilyl)imidazole in a chlorinated hydrocarbon solvent, such as, methylene chloride.

[0036] A compound of formula XXIV is converted to the corresponding compound of formula Id wherein R is fluorine by reaction with [3S-(3 α ,5 β ,Z)]-2-[2-methylene-3-fluoro-5-[[[(1,1-dimethylethyl)dimethylsilyl]oxy]cyclohexylidene]ethyl]diphenylphosphine oxide in an ether like solvent such as tetrahydrofuran in the presence of a base such as n-butyllithium.

[0037] The compounds of formula I can be administered orally, for example, in the form of tablets, coated tablets, dragees, hard or soft gelatin capsules, solutions, emulsions or suspensions. The administration can, however, also be effected rectally, for example, in the form of suppositories or parenterally, for example, in the form of injection solutions.

[0038] A composition in accordance with the invention can be processed with pharmaceutically inert, inorganic or organic excipients for the manufacture of tablets, coated tablets, dragees and hard gelatin capsules. Lactose, corn starch or derivatives thereof, talc, stearic acid or its salts, can be used as such excipients, for example, for tablets, dragees and hard gelatin capsules. Suitable excipients for soft gelatin capsules are, for example, vegetable oils, waxes, fats, semi-solid and liquid polyols; depending on the nature of the active ingredient. No excipients are, however, usually required in the case of soft gelatin capsules. Suitable excipients for the preparation of solutions and syrups, are, for example, water, polyols, saccharose, invert sugar, glucose.

[0039] Suitable excipients for injection solutions are, for example, water, alcohols, polyols, glycerol, vegetable oils.

[0040] Suitable excipients for suppositories are, for example, natural or hardened oils, waxes, fats, semi-liquid or liquid polyols.

[0041] Moreover, the pharmaceutical preparations can contain preservatives, solubilizers, stabilizers, wetting agents, emulsifiers, sweeteners, colorants, flavorants, salts for varying the osmotic pressure, buffers, masking agents or antioxidants.

[0042] The compounds of formula I as described above can be administered orally or by injection, for the treatment of neoplastic diseases such as leukemia, to warmblooded animals which need such treatment. More specifically, the compounds of formula I as described above can be administered orally to an adult human in dosages that are in the range of 0.25 to 50 μ g per day for the treatment of neoplastic diseases such as leukemia or breast cancer.

[0043] The compounds of formula I as described above can be administered orally, for the treatment of hyperproliferative skin diseases such as psoriasis, basal cell carcinomas, disorders of keratinization, and keratosis, to warm-blooded animals which need such treatment. More specifically, the compounds of formula I as described above can be administered orally to an adult human in dosages that are in the range of 0.25 to 50 μ g per day for the treatment of hyperproliferative skin diseases such as psoriasis, basal cell carcinomas, disorders of keratinization, and keratosis. These compounds can be administered orally for the treatment of acne in humans at a dosage of 0.25 to 50 μ g per day; preferably 0.5 to 5 μ g per day.

[0044] The compounds of formula I as described above can be administered topically, for the treatment of hyperproliferative skin diseases such as psoriasis, basal cell carcinomas, disorders of keratinization, and keratosis, to warm-blooded animals which need such treatment. More specifically, the compounds of formula I as described above can be administered topically in dosages that are in the range of 0.5 to 100 μ g per gram of topical formulation per day, for the treatment of hyperproliferative skin diseases such as psoriasis, basal cell carcinomas, disorders of keratinization, and keratosis.

[0045] The compounds of formula I as described above can also be administered topically for the treatment of sebaceous gland diseases such as acne or seborrheic dermatitis.

[0046] The useful activity of compounds of formula I as agents for the treatment of neoplastic diseases can be demonstrated by the following test procedures.

HL-60 Cell Differentiation

[0047] The induction of differentiation of HL-60 cells was assayed by measuring their oxidative burst potential via

the reduction of nitrobluetetrazolium (NBT).

[0048] HL-60 cells were maintained in RPMI 1640 medium supplemented with 10% fetal calf serum (FCS), 2mM L-glutamine, 1mM sodium pyruvate, 1% non-essential amino acids, 50 U/ml penicillin and 50 µg/ml streptomycin (RPMI/FCS). HL-60 cells (30,000 cells/90 µl of RPMI/supplemented medium) were seeded into flat-bottomed microtiter wells. Immediately after seeding, 10 µl of test compounds diluted in supplemented RPMI medium were added to the wells at the same time to yield final concentrations between 10^{-11} and 10^{-6} M (starting from stock solutions of 10^{-2} M in ethanol stored at -20 C and protected from light). After 3 days, the medium was removed from the wells with a multichannel pipette and replaced with 100 µl of NBT solution (1 mg/ml in phosphate buffered saline (PBS) with 200 nM phorbol myristate acetate (PMA). Following an additional hour incubation at 37 C the NBT solution was removed and 100 µl of 10% sodium dodecyl sulfate (SDS) in 0.01 N HCl was added. The amount of the reduced NBT was quantified photometrically at 540 nm using an automated plate reader. The mean of 3 wells was calculated. S.E.M. were between 5 and 10%. Values were expressed as percent of maximal differentiation achieved with 100-1000 nM calcitriol in the same experiment. The concentration (nM) leading to 50% of this maximal value is determined graphically and given in Table I as ED₅₀.

TABLE I

COMPOUND	ED ₅₀ (nM)
1,25-Dihydroxy-cholecalciferol	6.0
1,25-Dihydroxy-16-ene-23-yne 26,27-hexafluoro-20-epi-cholecalciferol	2.0
1,25-Dihydroxy-16,23E-diene-26, 27-hexafluoro-20-epi-cholecalciferol	0.37
1,25-Dihydroxy-16,23Z-diene-26, 27-hexafluoro-20-epi-cholecalciferol	2.3
1,25-Dihydroxy-16-ene-23-yne-20-epi-cholecalciferol	15.0
1,25-Dihydroxy-16-ene-20-epi-cholecalciferol	2.5
1α-Fluoro-25-hydroxy-16,23Z diene-26,27-hexafluoro-20-epi-cholecalciferol	20.0
1,25-Dihydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-19-nor-cholecalciferol	4.5
The 1,25-dihydroxy derivatives in Table I are not according to the invention.	

Antiproliferative Activity in T47-D and MCF-7 Breast Carcinoma Cells

[0049] The two cell lines used in these experiments and their growth requirements are listed below:

1. T47-D breast carcinoma cells were grown in RPMI 1640 medium supplemented with 10 µg/ml bovine insulin, and 10% fetal bovine.
2. MCF-7 breast carcinoma cells were grown in MEM (Eagles) supplemented with non-essential amino acids, 1 mM sodium pyruvate, 10 µg/ml bovine insulin, and 10% fetal bovine serum.

[0050] Cells were grown in appropriate medium to late log phase (~80% confluency). T47-D or MCF-7 cells are then trypsinized and seeded at 4000 or 2000 cells/well, respectively. At 24 hour post seeding, serial dilution of ethanol-solubilized drugs are prepared in the same medium and added to triplicate wells at a final concentration of 1,000 to 0.1 nM and 0.1% ethanol. On day 3 to 7 post drug addition, 50 µl of a 5 mg/ml MTT solution (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide in phosphate-buffered saline) is added to each well and incubation is continued at 37 C for 2.5 hours. The plates are then spun briefly by centrifugation at 800 xg for 5 minutes, medium is aspirated from wells, and 50 µl ethanol/well is added to dissolve the formazan formed during the incubation period with MTT. After a 15 minute shaking, the optical density is determined for each well in an automatic plate reader at 570 and 660 nm. Percent inhibition of cell growth is calculated by comparing optical densities of cells treated with test compounds to those of cells treated only with 0.1% ethanol. IC₅₀ values are determined based on the Reed and Muench formula (Reed, L.J., and H. Muench, A simple method of estimating fifty percent endpoint. Am. J. Hyg. 27:493-497 (1938)) and the results are presented below in Table II.

TABLE II

COMPOUND	T47-D	MCF-7
1,25-Dihydroxy-cholecalciferol	81.0	149
1,25-Dihydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol	6.7	0.95
1,25-Dihydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol	10.0	ND
1 α -Fluoro-25-hydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol		11
1 α -Fluoro-25-hydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol		9
1,25-Dihydroxy-16-ene-23-yne-26,27-bishomo-20-epi-cholecalciferol		11
1,25-Dihydroxy-16-ene-23-yne-26,27-bishomo-19-nor-20-epi-cholecalciferol		12
1 α -Fluoro-25-hydroxy-16,23E-diene-26,27-bishomo-20-epi-cholecalciferol		30
The 1,25-dihydroxy derivatives in Table II are not according to the invention.		

[0051] The useful activity of compounds of formula I as agents for the treatment of hyperproliferative skin disease can be demonstrated by the following.

Inhibition of Keratinocytes Proliferation

[0052] HaCaT cell line - The immortalized human cell line HaCaT was used. ³H-thymidine incorporation was measured in exponentially growing cultures after 6 days of culture in presence of the test compound.

[0053] Cell culture - HaCaT cells were cultured in Dulbecco's Modified Eagle Medium (DMEM) and Nutrient Mixture Ham's F12 (F12), 3:1 (v/v, ICN) containing 4.5 g/l glucose and supplemented with 10% fetal calf serum (Gibco, FCS), L-glutamine (Gibco, 2mM), penicillin (Gibco, 50 IU/ml), streptomycin (Gibco, 50 µg/ml), EGF (10 ng/ml), hydrocortisone (400 ng/ml), cholera toxin (8.5 ng/ml) and insulin (5 ng/ml). The cells were maintained in a humidified atmosphere containing 5% CO₂ and 95% air and passaged every 3-4 days.

[0054] Inhibition of ³H-thymidine uptake - HaCaT cells (250 cells in complete culture medium) were seeded into 96-well culture dishes and incubated at 37 °C with 5% CO₂ and 95% air for 6 days. Inhibitors, dissolved at 10 x concentration in 1% ethanol, were added immediately at the beginning of the assay. ³H-thymidine (5 Ci/mmol, Amersham) was added at a concentration of 1 µCi/well and cells were pulse-labeled for the last 6 hours of the growth period. Cells were then trypsinized for 10 minutes at 37 °C under a vigorous agitation and harvested on to a 96-well GF/C filter plate (Uni Filter, Packard) using a Micro Mate 196 cell harvester (Packard). After drying at 40 °C under vacuum for 20-30 minutes, 20 µl of Micro Scint 0 scintillator (Packard) were added and the radioactivity bound to the filters was counted on a TOP COUNT (Packard). The results are set forth in Table III.

TABLE III

COMPOUND	IC ₅₀ (nM)
1,25-Dihydroxy-cholecalciferol	55.0
1,25-Dihydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol	50.0
1,25-Dihydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol	0.7
1,25-Dihydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-cholecalciferol	ND
1,25-Dihydroxy-16-ene-23-yne-20-epi-cholecalciferol	93.0
1,25-Dihydroxy-16-ene-20-epi-cholecalciferol	4.0
The 1,25-dihydroxy derivatives in Table III are not according to the invention.	

[0055] The useful activity of compounds of formula I as agents for the treatment of sebaceous gland diseases can be demonstrated by the following.

Inhibition of Human Sebocyte Proliferation In Vitro

[0056] Sebaceous cells were isolated from adult human sebaceous glands by a combination of enzymatic and me-

chanical methods (Doran et al., Characterization Of Human Sebaceous Cells In Vitro, J. Invest. Dermatol. 96:341-8 (1991)). The cells were cultured in Iscove's medium containing 10% fetal calf serum and 4 µg/ml dexamethasone on a layer of growth-arrested 3T3 mouse fibroblasts. Cells were plated in medium without the test compound and then given test compound in fresh medium 24-48 hours after the initial plating. The cultures were given fresh medium, containing the test compound, every 48 hours. On the day of harvesting, the cultures were rinsed with 0.03% EDTA (ethylenediamine tetraacetic acid) in PBS (phosphate-buffered saline), to remove only the 3T3 fibroblasts, followed by incubation in 0.05% trypsin/0.03% EDTA. The cells were suspended, mixed vigorously to prepare a single cell suspension and counted in a hemocytometer.

[0057] Stock solutions of compounds were made up as 10^{-2} M solutions in degassed 100% ethanol and stored at -20 °C in the dark. During experimental use, the solutions, which have been aliquoted, were brought to room temperature and used by diluting directly into complete medium to the appropriate concentration.

[0058] The compounds were tested for the inhibition of proliferation of sebaceous cell growth *in vitro* at 10^{-6} , 10^{-7} and 10^{-8} M.

The results are summarized in Table IV as the amount of compound necessary to inhibit the proliferation of sebaceous cells by 50% (ED_{50}) in nM as compared to a vehicle-treated culture.

TABLE IV

COMPOUND	ED_{50} (nM)
1,25-Dihydroxycholecalciferol	50
1,25-Dihydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol	<10
1,25-Dihydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol	<10
1,25-Dihydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-cholecalciferol	ND
1,25-Dihydroxy-16-ene-23-yne-20-epi-cholecalciferol	50
1,25-Dihydroxy-16-ene-20-epi-cholecalciferol	<1
The 1,25-dihydroxy derivatives in Table IV are not according to the invention.	

Calcium tolerance test in mice

[0059] Profound changes in calcium homeostasis strongly affect the weight development of mice.

Mice (25-30 g body weight) received daily subcutaneous administrations of the compound for 4 consecutive days. Body weight was registered just before and at the end of a 5 day treatment period. The "highest tolerated dose" (HTD) is the dose which results in zero weight gain during this treatment period. The results are set forth in Table V.

TABLE V

COMPOUND	HTD (µg/kg)
1,25-Dihydroxycholecalciferol	0.5
1,25-Dihydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol	6.0
1,25-Dihydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol	2.5
1,25-Dihydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-cholecalciferol	2.5
1,25-Dihydroxy-16-ene-23-yne-20-epi-cholecalciferol	100
1,25-Dihydroxy-16-ene-20-epi-cholecalciferol	30
1α-Fluoro-25-hydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-cholecalciferol	8.0
1,25-Dihydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-19-nor-cholecalciferol	3.0
The 1,25-dihydroxy derivatives in Table V are not according to the invention.	

[0060] The compounds of formula I when applied topically to the skin, reverse the condition associated with photodamage so as to moderate and retard the damage to the skin caused by sun exposure. The damage caused sun exposure may include premature aging, elastosis and wrinkling. This damage is more pronounced in older patients. By applying the compounds of formula I topically to the skin in an amount effective to reverse the conditions associated with photodamage, the acceleration of skin repair is accomplished to enhance the skin with a smoother and younger

appearance. The compounds of formula I should be applied to that portion or area of the skin which is affected by photodamage or in which treatment is desired. The use of the compounds of formula I in accordance with this invention can provide the effects of anti-aging and anti-wrinkling, as well as enhance the repair of sun damaged skin.

[0061] A compound of formula I, or a combination of compounds of formula I can be applied in accordance with this invention to human skin in conventional topical compositions. These compositions can be utilized to apply compounds of formula I to the skin of the body, particularly the face, legs, arms and hands. The preferred method of application of compounds of formula I topically to produce the best effects should start where a patient is between 30 and 55 years of age, when elastosis begins to appear and becomes more pronounced. Thereafter, this composition can be continuously applied to patients to reduce the effects and injury associated with sun exposure. Generally, it is preferred to begin the treatment when the patient reaches approximately 30 years of age and to continue the treatment throughout his life, in order that the effects of elastosis be reduced and to prevent any further progression of photodamage.

[0062] The compounds of formula I can be administered in accordance with this invention in any conventional suitable topical preparation, that is, in combination with any suitable conventional carrier useful for topical administration. Therefore, compounds of formula I can be administered in accordance with this invention in any suitable topical composition such as a cream, ointment, soap, solution, lotion, emulsion, shampoo. Generally, for most efficacious results, these topical compositions contain from .00001% to .1% by weight of the total composition of a compound of formula I, with amounts of from .0001% to .01% by weight of the composition being especially preferred. If desired, higher concentrations may be utilized depending upon the nature and extent of elastosis.

[0063] In formulating these compositions, any conventional non-toxic, dermatologically acceptable base or carrier in which a compound of formula I is stable can be utilized. The preferred compositions for use in this invention are the conventionally cosmetic compositions which can contain a cosmetically active ingredient which is topically administered to human skin to provide a cosmetic effect. Among the conventional cosmetically active materials which can be utilized in this composition are included: sunscreens, penetration enhancers, moisturizers, surfactants, emollient, colorants, conditioners, bacteriocides, astringents, detergents. The topical compositions of this invention can, if desired, contain suitable sunscreen agents. Any conventional sunscreen agent can be utilized in formulating the formulations containing compounds of formula I which can be utilized in accordance with this invention.

[0064] These topical compositions which contain compounds of formula I can contain any of the conventional excipients and additives commonly used in preparing topical compositions. Among the conventional additives or excipients, which can be utilized in preparing these cosmetic compositions in accordance with this invention are preservatives, thickeners, perfumes. In addition, the conventional antioxidants, such as butylated hydroxyanisoles (BHA), ascorbyl palmitate, propyl gallate, citric acid, butylated hydroxy toluene (BHT), ethoxyquin, tocopherol can be incorporated into these compositions. These topical compositions can contain conventional acceptable carriers for topical applications which are generally utilized in these compositions. These compositions may contain thickening agents, humectants, emulsifying agents and viscosity stabilizers, such as those generally utilized. In addition, these compositions can contain flavoring agents, colorants, and perfume which are conventional in preparing cosmetic compositions.

[0065] The topical compositions containing compounds of formula I can be applied to the skin and should be preferably applied once daily to the skin. For obtaining the reversal of the elastosis so as to impart to the skin a smooth and younger appearance, the topical compositions should be preferably applied for a period of 6 months. After that, compositions which contain compounds of formula I should be applied continually to maintain the effect of younger and smoother skin. These preparations can be applied according to the need of the patient as determined by the prescribing physician. In any event, the particular regimen for application of this composition to a patient will typically depend on the age, weight and skin condition of the individual.

[0066] The useful activity of compounds of formula I for reversing the photodamage in sun-exposed skin can be demonstrated as follows:

Repair of UVB-Induced Dermal Damage in the Hairless Mouse by Compounds of Formula I

[0067] Hairless mice (female, HRS/J strain, Jackson Labs, 5-7 weeks old at the start of the experiments) were housed in yellow light and irradiated three times per week with a bank of 8 Westinghouse Sunlamps (FS72T12 HO, peak irradiance at 313 nm) placed about 20 cm above the animals. UVB output was measured on an International Light Research Radiometer, model IL 1700, using a SEE240 detector. With this setup, the lamp output was approximately 3.5 mW/cm² and the time of exposure for 0.06 J/cm² was about 17 seconds; 1 MED is approximately 0.03 J/cm². The precise dose was delivered by a IL 844A Phototherapy Exposure Control. Daily doses were 0.03 J/cm² for two weeks, 0.06 J/cm² for two weeks and 0.08 J/cm² thereafter, until a total dose of approximately 4 J/cm² was accumulated. To effect repair of the dermal damage, the UVB irradiation was discontinued and the animals were divided into groups of approximately eight and treated three times per week with various concentrations of the vitamin D analogs dissolved in ethanol. All dosing was done under yellow light. A control group treated with acetone alone was included. Two-cm strips of dorsal skin were taken longitudinally down the center of the irradiated (and treated) area. Elastin fibers were

stained with Luna's aldehyde fuchsin and collagen by Van Gieson. In this model, repair is defined by the appearance of a normalized dermis extending from the epidermis down to the layer of compressed elastin. The extent of repair is reflected by the width of this zone. In these studies, since the width of the zone varies considerably, the area of the zone on a standard length of histological section is measured by image analysis. Compounds are tested at three doses and an approximate ED₅₀ calculated.

[0068] The results are given in Table VI.

TABLE VI

Compound	Dose (μg)	Repair Zone mm ²
Control		0.0028 0.0008
Compound A	0.1	0.0282 0.0092***
	0.5	0.0102 0.0067*

* P < 0.05, **P < 0.01, ***P < 0.001 vs vehicle control

[0069] Compound A is 1,25-dihydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol (not according to the invention).

[0070] The following Examples are provided to further describe the invention and are not intended to limit it in any way.

Example 1

[0071] To a magnetically stirred solution of 14.06 g (25 mmole) of [1R-[1α(S*),3αα,4β,7aβ]]-octahydro-4-hydroxy-β,7a-dimethyl-1H-indene-1-ethanol, in 75 ml of dichloromethane under an argon atmosphere was added 47 ml. (33.8 g, 331 mmole) of acetic anhydride, followed by 53.5 ml of pyridine and 1.0 g of dimethylaminopyridine. After 3.5 hr, 25 ml of methanol was added. After 15 min., the reaction mixture was poured into 750 ml of 1 M phosphoric acid, and 400 ml of dichloromethane was added. The phases were separated and the aqueous phase was extracted 2 more times with 500 ml of dichloromethane. The organic phases were washed with 250 ml of water followed by 750 ml of 1 M sodium bicarbonate to afford, after drying with Na₂SO₄, filtration, and evaporation, 19.85 g (quantitative yield) of [1R-[1α(S*),3αα,4β,7aβ]]-4-(acetyloxy)-octahydro-β,7a-dimethyl-1H-indene-1-ethanol acetate as a white solid.

Example 2

[0072] To a magnetically stirred solution of 20.81 g (20.6 mmole) of [1R-[1α(S*),3αα,4β,7a,β]]-4-(acetyloxy)-octahydro-β,7a-dimethyl-1H-indene-1-ethanol acetate in 200 ml of methanol under an argon atmosphere was added 8.18 g (77.2 mmole) of sodium carbonate. The suspension was stirred for 16 hr and then most of the methanol was removed under reduced pressure on a rotary evaporator. The concentrate was distributed between 300 ml of water and 250 ml of ether. The aqueous phase was extracted 3 more times with 250 ml of ether and the ether phases were washed with 300 ml of water, dried with Na₂SO₄, filtered, and evaporated to give 18.02 g of crude product. Chromatography on medium pressure LC (Waters 500) gave, on elution with 2:1 hexane-ethyl acetate, 1.42 g unhydrolyzed diacetate, 16.08 g (90% yield) of [1R-[1α(S*),3αα,4β,7aβ]]-4-(acetyloxy)-octahydro-β,7a-dimethyl-1H-indene-1-ethanol as a colorless oil.

Example 3

[0073] A two-liter three-neck flask fitted with magnetic stirring, thermometer, and dropping funnel and maintained under an argon atmosphere was charged with 6.62 ml (75.8 mmole) of oxalyl chloride in 50 ml of dichloromethane. The flask was cooled to -65 °C in a dry ice-acetone bath; then a solution of 10.7 ml (151 mmole) of dimethylsulfoxide in 125 ml of dichloromethane was added dropwise at a rapid rate over 20 min while maintaining the temperature at -62 to -63 °C. After the addition, the reaction was stirred for an additional 5 min.; then a solution of 16.0 g of [1R-[1α(S*),3αα,4β,7aβ]]-4-(acetyloxy)-octahydro-β,7a-dimethyl-1H-indene-1-ethanol in 200 ml of dichloromethane (dried over 4 Å molecular sieves) was added dropwise over 20 min. maintaining the temperature at -65 °C. During the addition a precipitate formed. After an additional 20 min at -70 °C, a solution of 42.1 ml (302 mmole) of triethylamine in 75 ml of dry dichloromethane was added over 15 min. The suspension was stirred for 45 min, the cooling bath was removed, and the reaction mixture allowed to reach room temperature (1.5 hr). Most of the dichloromethane was removed on a rotary evaporator under reduced pressure (room temperature bath) and the residue was equilibrated with 500 ml of water and 750 ml of ether. The aqueous phase was extracted 3 times with 750 ml of ether and the ether phases washed in a counter current manner with 500 ml of water to afford, after drying with Na₂SO₄, filtration, and evaporation under

reduced pressure, 15.36 g of crude product. Medium pressure chromatography on silica gel using 4:1 hexanes-ether as the eluent (fractions collected under a nitrogen atmosphere) gave 14.95 g (94% yield) of [1R-[1 α (S*),3 $\alpha\alpha$,4 β ,7 $\alpha\beta$]]-4-(acetyloxy)-octahydro- α ,7 α -dimethyl-1H-indene-1-acetaldehyde as a colorless oil.

Example 4

[0074] To a magnetically stirred solution of 2.0 g (7.95 mmole) of [1R [1 α (S*),3 $\alpha\alpha$,4 β ,7 $\alpha\beta$]]-4-(acetyloxy)-octahydro- α ,7 α -dimethyl-1H-indene-1-acetaldehyde in 10 ml of ether under an argon atmosphere was added 100 mg of 10% palladium on charcoal. After 20 min. at ambient temperature, the suspension was filtered. The filtrate was evaporated under reduced pressure. To the residue was added 1.40 g (9.5 mmole) of benzalacetone and 200 mg of 10% palladium on charcoal. The suspension was degassed by evacuating the flask and refilling with argon. Then the flask was partially immersed in a 230 °C oil bath for 30 min. After cooling, the contents of the flask were flash chromatographed on 100 g of silica gel to remove the more polar benzalacetone and its reduction product (benzylacetone) from the equilibrium mixture of Δ^{17E} , Δ^{17Z} , Δ^{16} , and Δ^{20} indene olefins, which are present in a ratio of 65:4:27:4, respectively. Medium pressure chromatography on silver nitrate impregnated silica gel columns (two passes) separated the desired product (Δ^{17E}) from the less polar (Δ^{17Z} , Δ^{16} , and Δ^{20} in order of elution) olefins. Purification could be followed by TLC plates which were sprayed with a 10% solution of silver nitrate in acetonitrile and air dried before use. Thus, a total of 976 mg (55% yield) of [3 α R-(1E,3 $\alpha\alpha$,4 β ,7 $\alpha\beta$)]-1-ethylideneoctahydro-7 α -methyl-1H-inden-4-ol acetate was obtained as a colorless oil.

[0075] The more polar olefin mixture (420 mg., 24% yield) could be conveniently reequilibrated by addition to a subsequent aldehyde fragmentation reaction.

Example 5

[3 α S-[3(1R*,2R*S*, 3 $\alpha\beta$,7 β ,7 $\alpha\alpha$)]-3 α ,4,5,6,7,7 α -Hexahydro-3-[2-hydroxy-1-methyl-4-(trimethylsilyl)-3-butynyl]-3 α -methyl-1H-inden-7-ol acetate.

[0076] A flame dried two-liter three-neck flask fitted with stirrer, dropping funnel, thermometer, and argon inlet was charged with 22.3 g (100 mmole) of [3 α R-(1E,3 $\alpha\alpha$,4 β ,7 $\alpha\beta$)]-1-ethylideneoctahydro-7 α -methyl-1H-inden-4-ol acetate and 300 ml of dry dichloromethane. The solution was cooled to -20 °C with an acetone bath by addition of dry ice as necessary. At this point, the reagents were added incrementally. Initially, 200 ml (200 mmole) of 1M dimethylaluminum chloride in hexane was added rapidly dropwise over 5 min followed, after 10 min, by slow dropwise addition over 1 hr of 50 ml of a solution of 55.5 g (500 mmole) of 3-trimethylsilylpropynal diluted to 260 ml with dry dichloromethane. The addition of reagents was repeated in this manner 4 more times using 100 ml (100 mmole) of 1M dimethylaluminum chloride in hexane added rapidly followed by 50 ml of a solution of 3-trimethylsilylpropynal added slowly over 1 hr. After the final addition, TLC (CH₂Cl₂-Et₂O, 98:2) confirmed the absence of starting olefin, and the reaction mixture was poured with vigorous stirring into 2 l of 20% Rochelle salt solution to which was added about 500 g of ice (final temperature was 20 °C). The mixture was made alkaline by addition of 25 ml of 4 N sodium hydroxide, and 1 l of ether was added. The phases were separated and the ether phase was washed with 500 ml of brine. The aqueous phases were extracted in a counter current manner two times with 1.5 l of ether to afford, after drying with Na₂SO₄, filtration and evaporation under reduced pressure, 78 g of an oil. Medium pressure chromatography on silica gel was complicated by the difficulty of separating the products from a by-product, 4-trimethylsilyl-3-butyn-2-ol. Subsequently, advantage was taken of the volatility of the 4-trimethylsilyl-3-butyn-2-ol to remove it from the chromatographic fractions. Thus, there was obtained 25.54 g of more polar pure major isomer, 2.07 g of a 87:13 mixture (estimated by NMR) of major and minor isomers, and 0.53 g of pure minor isomer. The total yield of products was 79% (77% yield of the major isomer).

[0077] An analytical sample of the major isomer, [3 α S-[3(1R*,2S*, 3 $\alpha\beta$,7 β ,7 $\alpha\alpha$)]-3 α ,4,5,6,7,7 α -hexahydro-3-[2-hydroxy-1-methyl-4-(trimethylsilyl)-3-butynyl]-3 α -methyl-1H-inden-7-ol acetate, obtained as a white solid from ethyl acetate-hexane, m.p. 66-67 °C.

[0078] An analytical sample of the minor isomer, [3 α S-[3(1R*,2R*, 3 $\alpha\beta$,7 β ,7 $\alpha\alpha$)]-3 α ,4,5,6,7,7 α -hexahydro-3-[2-hydroxy-1-methyl-4-(tri-methylsilyl)-3-butynyl]-3 α -methyl-1H-inden-7-ol acetate, obtained as a colorless oil.

Example 6

[0079] To a magnetically stirred solution of 24.2 g (69 mmole) of [3 α S-[3(1R*,2S*,3 $\alpha\beta$,7 β ,7 $\alpha\alpha$)]-3 α ,4,5,6,7,7 α -hexahydro-3-[2-hydroxy-1-methyl-4-(trimethylsilyl)-3-butynyl]-3 α -methyl-1H-inden-7-ol acetate in 300 ml of dry dichloromethane and 23.3 ml of dry pyridine was added 25.0 g (145 mmole) of phenyl chloro-thionoformate. A yellow amber color developed. The reaction mixture was stirred for 3 hr and then 10 ml of methanol was added. After 15 min, the reaction mixture was transferred to a separatory funnel containing 1 liter of ether. The ether phase was washed suc-

cessively with 2 x 250 ml of 1 M phosphoric acid, 500 ml of water, and 500 ml of 1 M sodium bicarbonate. The aqueous phases were extracted in a counter current manner with 500 ml of ether. The combined ether phases were dried with Na₂SO₄, filtered, and evaporated under reduced pressure to give 47 g of crude product. Flash chromatography on 500 g of silica gel afforded 33.4 g (quantitative yield) of the thionocarbonate ester.

[0080] To a two-liter three-neck flask fitted with stirrer, dropping funnel, thermometer, and argon inlet charged with 33.4 g (69 mmole) of the above thionocarbonate ester in 500 ml of toluene was added rapidly dropwise, over 5 min 135 ml (500 mmole) of tri-n-butyltin hydride, followed by 276 ml (276 mmole) of triethylborane over 15 min. After 1.3 hr, an additional 15 ml (56 mmole) of tri-n-butyltin hydride was added. After 50 min, the reaction mixture was poured into 500 ml of 10% sodium bicarbonate and 500 ml of ether was added. The aqueous phase was extracted two times with 500 ml of ether. The ether phases were washed in a countercurrent manner three times with 250 ml of water; and after drying with Na₂SO₄, filtration, and evaporation under reduced pressure, gave the crude product which was flash chromatographed on 500 g of silica gel. The product containing fractions in dichloromethane solution were washed with 10% sodium carbonate to remove the phenol formed in the reaction. Medium pressure chromatography required several passes to remove the by products and afforded 16.13 g (70%) of pure [3aS-[(1S*,3aβ,7β,7aα)]-3a,4,5,6,7,7a-hexahydro-3-[1-methyl-4-(trimethylsilyl)-3-butynyl]-3a-methyl-1H-inden-7-ol acetate as a colorless oil.

Example 7

[3aS-[(1S*),3aβ,7β,7aα)]-3a,4,5,6,7,7a-Hexahydro-3-[1-methyl-3-butynyl]-3a-methyl-1H-inden-7-ol.

[0081] To a solution of 5.01 g (15 mmole) of [3aS-[(1S*,3aβ,7β,7aα)]-3a,4,5,6,7,7a-hexahydro-3-[1-methyl-4-(trimethylsilyl)-3-butynyl]-3a-methyl-1H-inden-7-ol acetate in 50 ml of ethanol was added 30 ml (60 mmole) of 2N sodium hydroxide and heated at 80 °C for four hours. It was then diluted with 500 ml water-brine (1:1), and thoroughly extracted with ethyl acetate. The combined extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (4:1) to give 3.14 g (95.4%) of the title compound.

Example 8

[3aS-[(1S*),3aβ,7β,7aα)]-3a,4,5,6,7,7a-Hexahydro-3-[1-methyl-3-butynyl]-3a-methyl-7-[(trimethylsilyl)oxy]-1H-indene.

[0082] To a solution of 3.14 g (14.4 mmole) of [3aS-[(1S*),3aβ,7β,7aα)]-3a,4,5,6,7,7a-hexahydro-3-[1-methyl-3-butynyl]-3a-methyl-1H-inden-7-ol in 30 ml of anhydrous methylene chloride was added 4.22 ml (28.8 mmole) of 1-(trimethylsilyl)imidazole, and the reaction mixture was stirred at room temperature for 1 hour. It was then quenched by addition of crushed ice and stirring for 15 minutes. After dilution with water and brine, it was thoroughly extracted with hexane. The combined extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-methylene chloride (40:1), to give 3.55 g (85%) of the title compound.

Example 9

[1(S*),3aR-(3aα,4β,7aβ)]-3a,4,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-hydroxy-3-hexynyl)-7a-methyl-4-[(trimethylsilyl)oxy]-3H-indene.

[0083] To a solution of 612 mg (2.10 mmole) of [3aS-[(1S*),3aβ,7β,7aα)]-3a,4,5,6,7,7a-hexahydro-3-[1-methyl-3-butynyl]-3a-methyl-7-[(trimethylsilyl)oxy]-1H-indene in 15 ml of anhydrous ether at -78 °C was added 1.6 ml (2.52 mmole) of 1.6 M n-butyllithium in hexane under an argon atmosphere. After stirring for 1 hour at -78 °C, 1.5 ml (21 mmole) of anhydrous acetone was added and the stirring was continued for 1 hour. The reaction was quenched with water-brine (1:1) and extracted thoroughly with ethyl acetate. The combined extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. Separation by FLASH chromatography with hexane-ethyl acetate (9:1) regenerated 166 mg (27%) of starting material, and gave 521 mg (71%) of the title compound.

Example 10

[1(S*),3aR-(3aα,4β,7aβ)]-3a,4,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-hydroxy-3-hexynyl)-7a-methyl-3H-inden-4-ol.

[0084] To a stirred solution of 521 mg (1.50 mmole) of [1(S*),3aR-(3aα,4β,7aβ)]-3a,4,5,6,7,7a-hexahydro-1-(1,5-dime-

thyl-5-hydroxy-3-hexynyl)-7a-methyl-4-[(trimethylsilyl)oxy]-3H-indene in 10 ml of anhydrous tetrahydrofuran was added 3 ml (3 mmole) of 1 M tetrabutylammonium fluoride. The mixture was stirred at room temperature for 1.5 hours, then diluted with water and brine and extracted thoroughly with ethyl acetate. The combined extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was crystalline. It was recrystallized from hexane to give 360 mg (87%) of the title compound, m.p. 115-116 C (from hexane).

Example 11

[1(S*),3aR-(3a α ,7a β)]-3,3a,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-hydroxy-3-hexynyl)-7a-methyl-4H-inden-4-one.

[0085] To a solution of 240 mg (0.868 mmole) of [1(S*),3aR-(3a α ,4 β ,7a β)] 3a,4,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-hydroxy-3-hexynyl)-7a-methyl-3H-inden-4-ol in 7 ml of anhydrous methylene chloride was added 980 mg (2.6 mmole) of pyridinium dichromate. This mixture was stirred at room temperature for 2 hours, then additional 500 mg (1.33 mmole) of pyridinium dichromate was added, and stirring was continued for 2.25 hours. After addition of 30 ml ether and stirring for 15 min, the mixture was filtered through Celite, and the Celite pad was washed with 3 x 50 ml of ethyl acetate. The combined filtrates were washed with 20 ml 1N HCl, water, 40 ml 2N potassium bicarbonate, and a mixture of water and brine 1:1. The aqueous layers were reextracted with 2 x 100 ml ethyl acetate. The combined organic layers were dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate 2:1, to give 214 mg (89%) of the title compound.

Example 12

[1(S*),3aR-(3a α ,7a β)]-3,3a,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-[(trimethylsilyl)oxy]-3-hexynyl)-7a-methyl-4H-inden-4-one

[0086] To a solution of 214 mg (0.78 mmole) of [1(S*),3aR-(3a α ,7a β)]-3,3a,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-hydroxy-3-hexynyl)-7a-methyl-4H-inden-4-one in 7 ml of anhydrous methylene chloride was added 750 mg (4.7 mmole) of 1-(trimethylsilyl)-imidazole. The mixture was stirred for 17 hours at room temperature in an argon atmosphere, and then quenched with 5 ml of water and stirred for 20 min. It was then extracted with 3 x 100 ml ethyl acetate. The extracts were combined and washed with 5 x 50 ml of water and brine mixture, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (12:1) to give 255 mg (94%) of the title compound.

Example 13

1,25-Dihydroxy-16-ene-23-yne-20-epi-cholecalciferol (not according to the invention)

[0087] To a stirred solution of 730 mg (1.25 mmole) of [3S-(1Z,3a,5 β)]-[2-[3,5-bis[[1,1-dimethylethyl]dimethylsilyl]oxy]-2-methylenecyclohexylidene]ethyl]diphenylphosphine oxide, a known compound, in 7 ml anhydrous tetrahydrofuran at -78 C was added 0.78 ml (1.25 mmole) of 1.6M n-butyllithium in hexane dropwise in an argon atmosphere. After stirring for 5 min, to the thus obtained red colored solution was added dropwise a solution of 255 mg (0.736 mmole) of [1(S*),3aR-(3a α ,7a β)]-3,3a,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-[(trimethylsilyl)oxy]-3-hexynyl)-7a-methyl-4H-inden-4-one in 4.5 ml anhydrous tetrahydrofuran over a 10 min period, and the reaction mixture was stirred at -78 C for 90 min. The reaction was then quenched by addition of 10 ml of a 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate and was allowed to warm up to room temperature. Additional 30 ml of Rochelle salt/potassium bicarbonate solutions were added and the reaction mixture was extracted with 3 x 100 ml ethyl acetate. The extracts were washed three times with brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate 40:1 to give 467 mg (89%) of the trisilylated intermediate. To the solution of the trisilylated intermediate (467 mg) in 4.5 ml anhydrous tetrahydrofuran was added 7 ml (7 mmole) of a 1M tetrabutylammonium fluoride in tetrahydrofuran, and this mixture was stirred at room temperature in argon atmosphere for 17 hours. The reaction was then quenched by addition of 5 ml water and stirring for 20 min. Tetrahydrofuran was then removed in vacuum, the residue was diluted with water and extracted with 3 x 100 ml ethyl acetate. The extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate 1:5 to give 250 mg (83%) of crystalline title compound. It was recrystallized from 1.5 ml tetrahydrofuran by addition of 7 ml methyl formate; m.p. 148-149 C.

Example 14

[1(S*),3aR-(3a α ,4 β ,7a β)]-3a,4,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-hydroxy-3Z-hexenyl)-7a-methyl-3H-inden-4-ol

- 5 **[0088]** A mixture of 1.02 g (3.6 mmole) of [1(S*),3aR-(3a α ,4 β ,7a β)]-3a,4,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-hydroxy-3-hexenyl)-7a-methyl-3H-inden-4-ol, 13.5 ml ethyl acetate, 34 ml hexane, 1.3 ml absolute ethanol, 0.067 ml quinoline and 225 mg Lindlar catalyst was hydrogenated at room temperature for 3.5 hours. The reaction mixture was filtered through a Celite pad, which was subsequently washed with ethyl acetate. The combined filtrates were washed with 1N HCl, water, 2N potassium bicarbonate, water and brine, dried over sodium sulfate and evaporated to dryness.
- 10 The crude product was purified by preparative HPLC using silica gel with hexane-ethyl acetate 3:2. It gave 954 mg (92.6%) of the title compound, m.p. 88-90 C (from CH₂Cl₂-hexane).

Example 15

- 15 [1(S*),3aR-(3a α ,4 β ,7a β)]-3a,4,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-hydroxy-hexanyl)-7a-methyl-3H-inden-4-ol

- [0089]** To a solution of 601 mg (2.16 mmole) of [1(S*),3aR-(3a α ,4 β ,7a β)]-3a,4,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-hydroxy-3Z-hexenyl)-7a-methyl-3H-inden-4-ol in 35 ml anhydrous methylene chloride was added 120 mg of 1,4-bis(diphenyl-phospheno)butane, 1,5-cyclooctadiene rhodium tetrafluoroborate and one drop of mercury and then hydrogenated using Paar apparatus at room temperature and 50 p.s.i. pressure for 1.25 hours. TLC (hexane-ethyl acetate, 3:2) showed that the reaction was complete. The reaction mixture was filtered through a Celite pad, which was then washed with ethyl acetate. The combined filtrates were evaporated to dryness. Purification was done by FLASH chromatography with hexane-ethyl acetate 3:2, and then by preparative HPLC using a silica gel column with hexane-ethyl acetate 3:2, 100 ml/min. It gave 550 mg (91%) of crystalline title compound

Example 16

[1(S*),3aR(3a α ,7a β)]-3,3a,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-hydroxyhexanyl)-7a-methyl-4-inden-4-one

- 30 **[0090]** To a solution of 270 mg (0.963 mmole) of [1(S*),3aR-(3a α ,4 β ,7a β)]-3a,4,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-hydroxy-hexanyl)-7a-methyl-3H-inden-4-ol in 7 ml of anhydrous methylene chloride was added 1.58 g (4.2 mmole) of pyridinium dichromate. The reaction mixture was stirred at room temperature for 4.75 hrs. It was then diluted with 25 ml of ether, stirred for additional 15 min, and filtered through a Celite pad. The pad was washed with 3 x 50 ml of ether. The combined filtrates were washed with 40 ml of 2N potassium bicarbonate and water-brine mixture. The aqueous layers were extracted with 2 x 90 ml of ethyl acetate. The combined organic layers were dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (2:1) to give 254 mg (95%) of the title compound.

Example 17

- 40 [1(S*),3aR(3a α ,7a β)]-3,3a,5,6,7,7a-Hexahydro-1-(1,5-dimethyl-5-[(trimethylsilyl)oxy]-hexanyl)-7a-methyl-4H-inden-4-one

- 45 **[0091]** To a solution of 254 mg (0.912 mmole) of [1(S*),3aR-(3a α ,7a β)]-3,3a,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-hydroxy-hexanyl)-7a-methyl-4H-inden-4-one in 6 ml of anhydrous methylene chloride was added 0.8 ml (5.45 mmole) of 1-(trimethylsilyl)-imidazole. The reaction mixture was stirred at room temperature under argon for 4.25 hrs. It was quenched then by addition of 5 ml of water and stirring for 20 min., and extracted with 3 x 90 ml of ethyl acetate. The extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (12:1) to give 308 mg (96%) of the title compound.

Example 18

1,25-Dihydroxy-16-ene-20-epi-cholecalciferol (not according to the invention).

- 55 **[0092]** To a stirred solution of 850 mg (1.46 mmole) of [3S-(1Z,3 α ,5 β)]-2-[3,5-bis[[[1,1-dimethylethyl]dimethylsilyl]oxy]-2-methylenecyclohexylidene]-ethyl]diphenylphosphine oxide in 9 ml of anhydrous tetrahydrofuran at -78 C was added 0.91 ml (1.46 mmole) of 1.6M n-butyllithium in hexane dropwise in an argon atmosphere. After stirring for 5 min, to thus obtained red colored solution was added dropwise a solution of 308 mg (0.878 mmole) of [1(S*),3aR-3a α ,7a β)]-

3,3a,5,6,7,7a-hexahydro-1-(1,5-dimethyl-5-[(trimethylsilyl)oxy]-hexanyl)-7a-methyl-4H-inden-4-one in 4.5 ml of anhydrous tetrahydrofuran over a 10 min period. The reaction mixture was stirred at -78 °C for 2 hours and then quenched by addition of 10 ml of 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate and by warming up to room temperature. After addition of 30 ml of Rochelle salt/potassium bicarbonate, the mixture was extracted with 3 x 100 ml of ethyl acetate. The combined extracts were washed with brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography with hexane-ethyl acetate 50:1 to give 541 mg of silylated intermediate.

To the solution of this intermediate in 5 ml of anhydrous tetrahydrofuran was added 5.5 ml (5.5 mmole) of 1 M tetrabutyl ammonium fluoride in tetrahydrofuran and the solution was stirred at room temperature under argon for 23 hours. The reaction was quenched with 5 ml of water and stirring for 20 min. After addition of 25 ml brine, it was extracted with 3 x 100 ml of ethyl acetate. The combined extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was first purified by FLASH chromatography with hexane-ethyl acetate (1:9), and then by HPLC (silica gel column) with ethyl acetate to give 296 mg (81%) of crystalline title compound, m.p. 124-125 °C (methyl formate).

Example 19

[3aR-[1(S*),3a α ,4 β ,7a β]]-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-4-[(trimethylsilyl)oxy]-3H-indene

[0093] In a 100 ml heart-shaped flask fitted with a gas inlet was placed a solution of 1 g (3.44 mmole) of [3aS-[1(S*),3a β ,7 β ,7a α]-3a,4,5,6,7,7a-hexahydro-3-[1-methyl-3-butynyl]-3a-methyl-7-[(trimethylsilyl)oxy]-1H-indene in 25 ml anhydrous tetrahydrofuran. After cooling at -78 °C, 3.2 ml (5.16 mmole) of 1.6M n-butyllithium in hexane was added dropwise over a 10 minute period, and the resulting mixture was stirred for 30 minutes. After adding 10 ml anhydrous tetrahydrofuran, a stream of hexafluoroacetone was bubbled into the mixture for one minute. After 1 hour and 15 minutes of stirring TLC showed that the reaction was about 70% complete. At this point the reaction was quenched by addition of 25 ml saturated brine, following by warming to room temperature. It was then extracted with hexane. The combined organic extracts were washed with brine, dried over sodium sulfate and evaporated to dryness. Purification of the crude product was done by FLASH chromatography with hexane-ethyl acetate (10:1). This gave 270 mg (27%) of starting material and 1.15 g (73%) of the title compound.

[0094] In the second experiment, to the solution of the starting acetylene 1.38 g (4.75 mmole) in 20 ml of anhydrous tetrahydrofuran at -78 °C in argon atmosphere was added dropwise over 10 min, 4.5 ml (7.2 mmole) of 1.6M n-butyllithium in hexane. After stirring for 30 min, hexafluoroacetone was bubbled at a slow rate for 2 min, stirred for 30 min, bubbled for an additional minute, stirred for 60 min, and finally bubbled for 30 sec and stirred for 30 minutes. At this time TLC showed the reaction complete. In this experiment the reaction was quenched with 2N Rochelle salt, and the purification by FLASH chromatography was with hexane-ethyl acetate (14:1). It gave 1.97 g (91 %) of the title compound.

Example 20

[3aR-[1(S*),3a α ,4 β ,7a β]]-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-3H-inden-4-ol

[0095] To a solution of 1.17 g (2.56 mmole) of [3aR-[1(S*),3a α ,4 β ,7a β]]-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-4-[(trimethylsilyl)oxy]-3H-indene in 20 ml anhydrous tetrahydrofuran was added 7.12 ml (7.12 mmole) of 1M tetrabutyl ammonium fluoride in tetrahydrofuran and stirred in argon atmosphere for 3.5 hours. The reaction mixture was then diluted with 250 ml of water-brine 1:1 mixture, and extracted thoroughly with ethyl acetate. The extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. Purification of the crude product was done by FLASH chromatography to give 946 mg (96%) of the title compound; m.p. 75-76 °C after recrystallization from hexane.

Example 21

[1(S*),3aR-(3a α ,7a β)]-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-4H-inden-4-one

[0096] To a solution of 284 mg (0.74 mmole) of [3aR-[1(S*),3a α ,4 β ,7a β]]-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-3H-inden-4-ol in 10 ml of anhydrous methylene

chloride was added in portions 2.15 g (5 mmole) of pyridinium dichromate and the reaction mixture was stirred at room temperature for 5 hours. After addition of 50 ml of ether and stirring for 15 minutes, the reaction mixture was filtered through Celite pad, which was subsequently washed with 3 x 50 ml of ethyl acetate. The combined filtrates were washed with 2N HCl, water, 2N potassium bicarbonate, water and brine, dried over sodium sulfate, and evaporated to dryness.

The crude product was purified by FLASH chromatography with hexane-ethyl acetate (3:1), to give 259 mg (91%) of the crystalline title compound.

Example 22

1,25-Dihydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol (not according to the invention)

[0097] To a stirred solution of 987 mg (1.69 mmole) of [3S-(1Z,3 α ,5 β)]-2-[3,5-bis[[[1,1-dimethylethyl]dimethylsilyl]oxy]-2-methylenecyclohexylidene]ethyl]diphenylphosphine oxide in 10 ml of anhydrous tetrahydrofuran at -78 °C was added 1.05 ml (1.69 mmole) of 1.6M n-butyllithium in hexane dropwise in an argon atmosphere. After stirring for 15 minutes, to thus obtained red colored solution was added dropwise a solution of 259 mg (0.677 mmole) of [1(S*), 3aR-(3a α ,7a β)-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-4H-inden-4-one in 5 ml anhydrous tetrahydrofuran, and thus obtained reaction mixture was stirred at -78 °C for 1 hour. The reaction was then quenched by addition of 2N Rochelle salt solution and by allowing to warm up to room temperature. After addition of saturated brine, it was extracted thoroughly with ethyl acetate. The combined extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography with hexane-ethyl acetate (9:1) to give 370 mg of intermediate disilyl ether.

To the solution of this intermediate (370 mg) in 10 ml of anhydrous tetrahydrofuran was added 3 ml (3 mmole) of a 1M tetrabutylammonium fluoride in tetrahydrofuran, and thus obtained reaction mixture was stirred at room temperature overnight. It was then diluted with water and brine and extracted thoroughly with 4 x 50 ml of ethyl acetate. The combined extracts were washed with water and brine, dried over sodium sulfate, and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (1:4) to give 249 mg (71%) of the title compound as a foam.

Example 23

[3aR-{1(S*),3a α ,4 β ,7a β }] -3a,4,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3E-hexenyl]-3H-inden-4-ol

[0098] To a under argon stirred suspension of 165 mg (4.34 mmole) lithium aluminum hydride in 15 ml anhydrous tetrahydrofuran, cooled in an ice-bath, was first added 235 mg (4.34 mmole) of solid sodium methoxide, and then dropwise 334 mg (0.87 mmole) of [3aR-1(S*),3a α ,4 β ,7a β]]-3a,4,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-3H-inden-4-ol dissolved in 10 ml of anhydrous tetrahydrofuran. The thus obtained reaction mixture was heated under reflux (80 °C) for 2.5 hours, when TLC indicated that the reaction was complete. The mixture was cooled in an ice-bath, and then quenched carefully with 1 ml of water and 1 ml 2N sodium hydroxide. After addition of 20 ml of ether, it was stirred 0.5 hour, 2.2 g magnesium sulfate was added, stirred for 0.5 hour, filtered and evaporated to dryness. It gave 335 mg (100%) of crystalline title compound (TLC pure).

Example 24

[1(S*),3aR-(3a α ,7a β)]-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3E-hexenyl]-4H-inden-4-one

[0099] To a solution of 256 mg (0.66 mmole) of [3aR-{1(S*),3a α ,4 β ,7a β }] -3a,4,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3E-hexenyl]-3H-inden-4-ol in 10 ml of anhydrous methylene chloride was added in portions of 1.12 g (3 mmole) of pyridinium dichromate, and the reaction mixture was stirred at room temperature for 4 hours. After addition of 25 ml of ether and stirring for 15 minutes, the reaction mixture was filtered through a Celite pad, which was subsequently washed with 3 x 20 ml of ethyl acetate. The combined filtrates were washed with 2N potassium bicarbonate, water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with methylene chloride-ethyl acetate (9:1) to give 230 mg (90%) of crystalline title compound.

Example 25

[1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-(trimethylsilyl)oxy-1-methyl-5-trifluoromethyl]-3E-hexenyl]-4H-inden-4-one

[0100] To a stirred solution of 268 mg (0.697 mmole) of [1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3E-hexenyl]-4H-inden-4-one in 8 ml of anhydrous methylene chloride at room temperature was added 0.92 ml (6.27 mmole) of 1-(trimethylsilyl)-imidazole, and the reaction mixture was stirred at room temperature overnight. The reaction mixture was then diluted with water and extracted thoroughly with hexane. The combined extracts were washed with water and brine to neutrality, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate 9:1 to give 310 mg (97%) of the title compound.

Example 26

1,25-Dihydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol (not according to the invention)

[0101] To a stirred solution of 725 mg (1.24 mmole) of [3S-(1Z,3 α ,5 β)]-[2-[3,5-bis(((1,1-dimethylethyl)dimethylsilyl)oxy)-2-methylene cyclohexylidene]ethyl]diphenylphosphine oxide in 15 ml of anhydrous tetrahydrofuran at -78 °C was added 0.78 ml (1.24 mmole) of 1.6M n-butyllithium in hexane dropwise in an argon atmosphere. After stirring for 5 min, to thus obtained red colored solution was added dropwise a solution of 307 mg (0.672 mmole) of [1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-(trimethylsilyl)oxy-1-methyl-5-(trifluoro-methyl)-3E-hexenyl]-4H-inden-4-one in 9 ml of anhydrous tetrahydrofuran. The reaction mixture was stirred at -78 °C for 2.5 hrs and then quenched with water and allowed to warm up to room temperature. After further dilution with water, it was extracted with 3 x 30 ml ethyl acetate. The combined extracts were washed with water and brine, dried over sodium sulfate and evaporated to dryness. Thus obtained disilyl ether intermediate was purified by FLASH chromatography with hexane-ethyl acetate 9:1 to give 449 mg of pure intermediate (the 25-silyloxy group was hydrolyzed during the reaction).

[0102] To the solution of this intermediate (449 mg) in 10 ml anhydrous tetrahydrofuran was added 3.6 ml (3.6 mmole) of a 1M tetrabutyl ammonium fluoride in tetrahydrofuran and thus obtained reaction mixture was stirred at room temperature overnight. The reaction was quenched with ice, stirred 15 min, diluted with water and brine, and extracted with ethyl acetate. The combined extracts were washed with water, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (1:2) to give the title compound as white foam, 291 mg (83%).

Example 27

[3aR-[1(S*),3a α ,4 β ,7a β]-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3Z-hexenyl]-3H-inden-4-ol

[0103] A mixture of 1.62 g (4.21 mmole) of [3aR-1(S*),3a α ,4 β ,7a β]-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-3H-inden-4-ol, 16 ml of ethyl acetate, 40 ml of hexane, 1.6 ml of absolute ethanol, 0.080 ml of quinoline and 320 mg of Lindlar catalyst was stirred under hydrogen atmosphere for 70 min at room temperature. It was then filtered over Celite and the filter cake was washed with 3 x 60 ml of ethyl acetate. The filtrate was washed with 25 ml of 1N HCl and 4 times with a mixture of water and brine. The aqueous layers were extracted with 2 x 90 ml of ethyl acetate. The combined organic layers were dried over sodium sulfate and evaporated to dryness. The crude product was purified by HPLC on silica gel column with hexane-ethyl acetate (3:1), flow 100 ml/min to give 1.51 g (93%) of the title compound as white solid.

Example 28

[1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3Z-hexenyl]-4H-inden-4-one

[0104] To a solution of 750 mg (1.94 mmole) of [3aR-[1(S*),3a α ,4 β ,7a β]-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3Z-hexenyl]-3H-inden-4-ol in 14 ml methylene chloride was added, in portions, 3.64 g (9.68 mmole) of pyridinium dichromate, and the reaction mixture was stirred at room temperature for 5 hours. To the reaction mixture was then added 30 ml of ether, stirred for 15 minutes, filtered through a Celite pad and the pad was washed with 3 x 50 ml of ether. The combined filtrates were washed with 2N potassium

bicarbonate (50 ml) and three times with water and brine 1:1. The aqueous layers were extracted with 2 x 100 ml of ethyl acetate. The combined organic layers were dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate to give 720 mg (96.5%) of the title compound as white solid.

Example 29

[1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-Hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-trimethylsilyloxy-1-methyl-5-trifluoromethyl-3Z-hexenyl]-4H-inden-4-one

[0105] To a stirred solution of 720 mg (1.87 mmole) of [1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-trifluoromethyl-3Z-hexenyl]-4H-inden-4-one in 13 ml of anhydrous methylene chloride at room temperature was added 1.86 ml (12.68 mmole) of 1-trimethylsilyl-imidazole, and the reaction mixture was stirred under argon atmosphere for 17 hours. The reaction was then quenched with 7 ml water and stirring for 15 min., 20 ml of brine was added and extracted with 3 x 100 ml of ethyl acetate. The combined extracts were washed five times with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (9:1) to give 825 mg (96.5%) of the amorphous title compound.

Example 30

1,25-Dihydroxy-16,23Z-diene-26,27-hexafluoro-20-*epi*-cholecalciferol (not according to the invention)

[0106] To a stirred solution of 555 mg (0.952 mmole) of [3S-(1Z,3 α ,5 β)]-[2-[3,5-bis[(1,1-dimethylethyl)dimethylsilyl]oxy]-2-methylenecyclohexylidene]ethyl]diphenylphosphine oxide in 6 ml of anhydrous tetrahydrofuran at -78 C was added 0.595 ml (0.952 mmole) of 1.6M n-butyllithium in hexane dropwise in an argon atmosphere. After stirring for 5 min, to the thus obtained red colored solution was added dropwise a solution of 270 mg (0.591 mmole) of [1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-trimethylsilyloxy-1-methyl-5-trifluoromethyl-3Z-hexenyl]-4H-inden-4-one in 4 ml of anhydrous tetrahydrofuran over a 10 min period. The reaction mixture was stirred at -78 C for 1.75 hours, and then was quenched by addition of 10 ml of 2N Rochelle salt and by warming up to room temperature. It was then diluted with 30 ml of 2N Rochelle salt and extracted with 3 x 100 ml of ethyl acetate. The combined extracts were washed three times with brine, dried over sodium sulfate and evaporated to dryness. The crude silylated intermediate was purified by FLASH chromatography with hexane-ethyl acetate (10:1) to give 355 mg of pure intermediate.

To the solution of this intermediate in 4 ml of anhydrous tetrahydrofuran was added 3.8 ml (3.8 mmole) of 1M tetrabutyl ammonium fluoride in tetrahydrofuran under argon, and the reaction mixture was stirred at room temperature for 19 hours. The reaction was then quenched with 5 ml water and stirring for 15 min. After dilution with 20 ml of brine, it was extracted with 3 x 90 ml of ethyl acetate. The combined extracts were washed 4 times with mixture of water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (1:3), to give 237 mg (77%) of crystalline title compound; m.p. 120-122 C (from tetrahydrofuran-methyl formate recrystallization).

Example 31

1,25-Dihydroxy-16,23Z-diene-26,27-hexafluoro-20-*epi*-19-norcholecalciferol (not according to the invention)

[0107] A solution of 700 mg (1.23 mmole) of [3R-(3 α ,5 β ,Z)-3,5-bis[(1,1-dimethylethyl)dimethylsilyl]oxy]cyclohexylidene]ethyl]diphenylphosphine oxide in 7 ml of anhydrous tetrahydrofuran was cooled in a dry ice bath to -78 C and treated with 0.77 ml (1.23 mmole) of 1.6M n-butyllithium in hexane and stirred for 5 min. under an argon atmosphere. To this was added dropwise in the course of 10 min. a solution of 277 mg (0.607 mmole) of [1(S*),3aR-(3a α ,7a β)-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-trimethylsilyloxy-1-methyl-5-trifluoromethyl-3Z-hexenyl]-4H-inden-4-one in 4 ml of anhydrous tetrahydrofuran. Thus obtained reaction mixture was stirred at -78 C for 2 hours. After addition of 10 ml of 2N Rochelle Salt, it was warmed up to room temperature. After further dilution with 25 ml of 2N Rochelle Salt, it was extracted with 3 x 100 ml of ethyl acetate. The organic layers were washed with brine, dried over sodium sulfate and evaporated to dryness. The residue was chromatographed on a silica gel column with hexane-ethyl acetate (10:1), followed by elution with hexane-ethyl acetate (1:3) to recover the ring A precursor. It gave 329 mg of the ring A disilylated title compound.

To the solution of 329 mg disilylated title compound in 2.5 ml of anhydrous tetrahydrofuran was added 4.4 ml (4.4 mmole) of 1 M tetrabutylammonium fluoride in tetrahydrofuran, and the reaction mixture was stirred at room temperature

for 65 hours. It was then quenched with 10 ml water, stirred for 15 min., 20 ml of brine was added and extracted with 3 x 90 ml of ethyl acetate. The combined organic layers were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was first purified by FLASH chromatography with hexane-ethyl acetate (1:8), and then by HPLC on a silica gel column, to give 210 mg (68%) of the title compound as white foam.

Example 32

1 α -Fluoro-25-hydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-cholecalciferol

[0108] To a solution of 460 mg (0.977 mmole) 5 β ,Z)-2-[2-[2-methylene-3-fluoro-5-[[[(1,1-dimethylethyl)dimethylsilyl]oxy] cyclohexylidene]ethyl]diphenyl phosphine oxide in 6 ml of anhydrous tetrahydrofuran at -78 C was added 0.61 ml (0.976 mmole) of 1.6M n-butyllithium in hexane dropwise in an argon atmosphere. After stirring for 5 min., a solution of 277 mg (0.607 mmole) of [1(S*),3aR-(3 α ,7 α β)-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-(trimethylsilyl)oxy-1-methyl-5-(trifluoromethyl)-3Z-hexenyl]-4H-inden-4-one in 4 ml of anhydrous tetrahydrofuran was added dropwise over a 10 min. period. The reaction mixture was then stirred at -78 C for 2 hrs. It was then quenched with 10 ml of 2N Rochelle Salt, and warmed up to room temperature. After a further dilution with 30 ml 2N Rochelle Salt, it was extracted with 3 x 100 ml ethyl acetate. The combined organic layers were washed with brine, dried over sodium sulfate and evaporated to dryness to give 630 mg of crude intermediate. It was purified by FLASH chromatography with hexane-ethyl acetate (10:1), to give 220 mg of disilylated title compound. (The 25-silyl ether is hydrolyzed during this procedure).

To the solution of 220 mg of disilylated intermediate in 4 ml of anhydrous tetrahydrofuran was added 2.5 ml (2.5 mmole) of a 1M tetrabutylammonium fluoride in tetrahydrofuran under argon, and then stirred at room temperature for 18 hrs. At that time, 5 ml of water was added and stirred 15 min, followed by addition of 20 ml of brine and extraction with 3 x 90 ml of ethyl acetate. The combined organic layers were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography with hexane-ethyl acetate (2:1) to give 170 mg (53%) of the title compound as white foam.

Example 33

1 α -Fluoro-25-hydroxy-16-ene-23-yne-26,27-hexafluoro-20-epicholecalciferol

[0109] To a solution of 740 mg (1.57 mmole) of [3S-(1Z,3 α ,5 β)]-2-[5-[[[(1,1-dimethylethyl)dimethylsilyl]oxy]-3-fluoro-2-methylenecyclohexylidene]ethyl]diphenylphosphine oxide in 8 ml of anhydrous tetrahydrofuran at -78 C was added with stirring 0.98 ml (1.57 mmole) of 1.6 M n-butyllithium in hexane, dropwise under argon. After stirring for five minutes, to thus formed red solution was added a solution of 262 mg (0.685 mmole) of [3aR-[1(S*),3 α ,7 α β]]-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-5-hydroxy-1-methyl-5-(trifluoromethyl)-3-hexynyl]-4H-inden-4-one in 5 ml tetrahydrofuran, dropwise over a 10 min. period. The reaction mixture was then stirred at -78 C for 2 hrs. It was then quenched by addition of 15 ml of a 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate, followed by warming to room temperature. After 30 ml of the same salts mixture was added, it was extracted 3 x 100 ml of ethyl acetate. The combined organic layers were washed three times with water and brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate (5:1) to give 250 mg of the monosilylated title compound.

To the solution of 250 ml silyl intermediate in 3 ml of anhydrous tetrahydrofuran was added 2.8 ml (2.8 mmole) of 1M tetrabutylammonium fluoride in tetrahydrofuran under argon. The reaction mixture was stirred at room temperature for 18 hrs. 10 ml of water was added and stirred for 15 min, then diluted with 20 ml brine and extracted with 3 x 90 ml ethylacetate. The organic layers were washed four times with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate (2:1) and by HPLC on a silica column. It gave 175 mg (49%) of amorphous title compound.

Example 34

1 α -Fluoro-25-hydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol

[0110] To a stirred solution of 350 mg (0.744 mmole) of [3S-(1Z,3 α ,5 β)]-2-[5-[[[(1,1-dimethylethyl)dimethylsilyl]oxy]-3-fluoro-2-methylenecyclohexylidene]-ethyl]diphenylphosphine oxide in 5 ml of anhydrous tetrahydrofuran at -78 C was added 0.463 ml (0.74 mmole) of 1.6 M n-butyl lithium in hexane dropwise under argon. After stirring for 5 min, to thus formed red solution was added 230 mg (0.503 mmole) of [3aR-[1(S*,3E),3 α ,7 α β]]-3,3a,5,6,7,7a-hexahydro-7a-methyl-1-[6,6,6-trifluoro-1-methyl-5-(trifluoro-methyl)-5-[(trimethylsilyl)oxy]-3-hexenyl]-4H-inden-4-one in 4.5 ml of an-

hydrous tetrahydrofuran dropwise over a 10 min period. The reaction mixture was stirred at -78 °C for two hours. It was then quenched by addition of 10 ml of a 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate and warmed up to room temperature. After addition of 30 ml of the same salt mixture, it was extracted with 3 x 100 ml of ethyl acetate. The combined organic layers were washed three times with water and brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate to give 236 mg of disilylated title compound.

To the solution of 236 mg disilyl intermediate in 3.5 ml anhydrous tetrahydrofuran was added 3 ml (3 mmole) of a 1M tetrabutylammonium fluoride in tetrahydrofuran under argon and the reaction mixture was stirred at room temperature for 18 hrs. Quenching was done by addition of 5 ml of water and stirring for 10 min. After dilution with 20 ml of brine, the mixture was extracted with 3 x 90 ml of ethyl acetate. The combined organic layers were washed with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate (2:1), and HPLC on a silica column, to give 140 mg (53%) of amorphous title compound.

Example 35

1,25-Dihydroxy-16-ene-23-yne-26,27-bishomo-20-epi-cholecalciferol (not according to the invention)

[0111] To a stirred solution of 435 mg (0.746 mmole) of [3S-(1Z,3α,5β)]-[2-[3,5-bis[[[(1,1-dimethylethyl)dimethylsilyl]oxy]-2-methylenecyclohexylidene]ethyl]diphenylphosphine oxide in 5 ml of anhydrous tetrahydrofuran at -78 °C was added 0.465 ml (0.744 mmole) of 1.6 M n-butyllithium in hexane dropwise under argon. After stirring for 5 min, to this obtained red solution was added a solution of 169 mg (0.451 mmole) of [3aR-[1(S*),3α,7aβ]]-1-[5-ethyl-1-methyl-5-[(trimethylsilyl)oxy]-3-heptynyl]-3,3a,5,6,7,7a-hexahydro-7a-methyl-4H-inden-4-one in 4 ml of anhydrous tetrahydrofuran, dropwise over a 10 min period. The reaction mixture was stirred at -78 °C for 2.5 hrs. It was then quenched by addition of 10 ml of a 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate and warmed up to room temperature. After 25 ml of the same salt mixture was added, it was extracted with 3 x 90 ml of ethyl acetate. The combined organic layers were washed three times with water and brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate (40:1) to give 260 mg of trisilylated title compound.

To the solution of 260 mg trisilylated intermediate in 3 ml of anhydrous tetrahydrofuran was added 3 ml (3 mmole) of 1M tetrabutylammonium fluoride in tetrahydrofuran in argon atmosphere. The reaction mixture was stirred at room temperature for 17 hrs, and then quenched by addition of 10 ml water and stirring for 10 min. After dilution with 20 ml water and brine, it was extracted with 3 x 80 ml of ethyl acetate. The combined organic layers were washed four times with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate (1:3), to give 158 mg (80%) of the title compound, which was crystallized from methyl formate, m.p. 103-105 °C.

Example 36

1,25-Dihydroxy-16-ene-23-yne-26,27-bishomo-19-nor-20-epi-cholecalciferol (not according to the invention)

[0112] To a stirred solution of 300 mg (0.525 mmole) of [3R-trans-[2-[3,5-bis[[[(1,1-dimethylethyl)dimethylsilyl]oxy]cyclohexylidene]ethyl]-diphenylphosphine oxide in 4 ml of anhydrous tetrahydrofuran at -78 °C was added 0.325 ml (0.520 mmole) of 1.6 M n-butyllithium in hexane, dropwise under argon. After stirring for 5 min, to this obtained red solution was added a solution of 77 mg (0.205 mmole) of [3aR-[1(S*),3α,7aβ]]-1-[5-ethyl-1-methyl-5-[(trimethylsilyl)oxy]-3-heptynyl]-3,3a,5,6,7,7a-hexahydro-7a-methyl-4H-inden-4-one in 2.5 ml of anhydrous tetrahydrofuran, dropwise over a 5 min period. The reaction was stirred at -78 °C for 2.5 hrs. and then quenched by addition of 10 ml of a 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate and warming up to room temperature. After a further dilution with 25 ml of the same salts mixture, it was extracted with 3 x 75 ml of ethyl acetate. The combined organic layers were washed three times with water and brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate to give 58 mg of trisilylated title compound.

To a solution of 124 mg of trisilyl intermediate (obtained from two preceding coupling reactions) in 2.5 ml of anhydrous tetrahydrofuran was added 2.5 ml (2.5 mmole) of 1M tetrabutylammonium fluoride in tetrahydrofuran under argon. The reaction mixture was stirred at room temperature for 40 hrs. It was then quenched by addition of 10 ml of water and stirring for 15 min. After dilution with 20 ml of water and brine, it was extracted with 3 x 70 ml of ethyl acetate. The combined organic layers were washed four times with water and brine, dried over sodium sulfate and evaporated to dryness. The crude product was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate

(1:7), to give 74 mg (25.3%) of the title compound, which was recrystallized from methyl formate, m.p. 130-132 C.

Example 37

1 α -Fluoro-25-hydroxy-16-ene-23-yne-26,27-bishomo-20-epi-cholecalciferol

[0113] To a stirred solution of 390 mg (0.829 mmole) of [3S-(1Z,3 α ,5 β)]-2-[5-[[[(1,1-dimethylethyl)dimethylsilyl]oxy]-3-fluoro-2-methylenecyclohexylidene]ethyl]diphenylphosphine oxide in 5 ml of anhydrous tetrahydrofuran at -78 C was added 0.53 ml (0.848 mmole) of 1.6 M n-butyllithium in hexane, dropwise under argon. After stirring for 5 min, to thus obtained red solution was added a solution of 200 mg (0.534 mmole) of [3aR-[1(S*),3 $\alpha\alpha$,7a β]]-1-[5-ethyl-1-methyl-5-[[trimethylsilyl]oxy]-3-heptynyl]-3,3a,5,6,7,7a-hexahydro-7a-methyl-4H-inden-4-one in 4.5 ml of anhydrous tetrahydrofuran, dropwise over a 10 min period. The reaction mixture was then stirred at -78 C for 2 hrs, and was quenched by addition of 10 ml of a 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate and warming up to room temperature. After dilution with 30 ml of the same salts mixture, it was extracted with 3 x 100 ml ethyl acetate. The combined organic layers were washed three times with water and brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography on silica gel column with hexane-ethyl acetate (30:1), to give 254 mg of disilylated title compound.

To a solution of 254 mg of disilyl intermediate in 3.5 ml anhydrous tetrahydrofuran was added under argon 3.5 ml (3.5 mmole) of 1M tetrabutylammonium fluoride in tetrahydrofuran. The reaction mixture was stirred at room temperature for 18 hrs, and then quenched by addition of 10 ml of water and stirring for 15 min. After dilution with 20 ml of water and brine, it was extracted with 3 x 80 ml of ethyl acetate. The combined organic layers were washed four times with water and brine, dried over sodium sulfate, and evaporated to dryness. The crude product was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate (7:4), and by HPLC on a silica column with hexane-ethyl acetate (3:2), to give 159 mg (67.6%) of amorphous title compound.

Example 38

1 α -Fluoro-25-hydroxy-16-23E-diene-26,27-bishomo-20-epi-cholecalciferol

[0114] To a stirred solution of 240 mg (0.51 mmole) of [3S-(1Z,3 α ,5 β)]-2-[5-[[[(1,1-dimethylethyl)dimethylsilyl]oxy]-3-fluoro-2-methylenecyclohexylidene]ethyl]diphenylphosphine oxide in 5 ml of anhydrous tetrahydrofuran at -78 C was added 0.319 ml (0.51 mmole) of 1.6 M n-butyllithium in hexane, dropwise under argon. After stirring for 5 min, to thus obtained red solution was added a solution of 103 mg (0.273 mmole) of [3aR-[1(S*,3E),3 $\alpha\alpha$,7a β]]-1-[5-ethyl-1-methyl-5-[[trimethylsilyl]oxy]-3-heptynyl]-3,3a,5,6,7,7a-hexahydro-7a-methyl-4H-inden-4-one in 4 ml of anhydrous tetrahydrofuran, dropwise over a 10 min period. The reaction mixture was stirred at -78 C for 2 hrs, then placed in freezer (-20 C) for one hour, quenched by addition of 10 ml of a 1:1 mixture of 2N Rochelle salt and 2N potassium bicarbonate and warmed up to room temperature. After dilution with additional 25 ml of the same salts mixture, it was extracted with 3 x 90 ml of ethyl acetate. The combined organic layers were washed three times with water and brine, dried over sodium sulfate and evaporated to dryness. The residue was purified by FLASH chromatography on a silica gel column with hexane-ethyl acetate (1:4), to give 145 mg of disilylated title compound. To a solution of 145 mg of disilyl intermediate in 3 ml anhydrous tetrahydrofuran was added 1.7 ml (1.7 mmole) of 1M tetrabutylammonium fluoride in tetrahydrofuran under argon. The reaction mixture was stirred at room temperature for 18 hrs, and then quenched by addition of 10 ml water and stirring for 15 min. It was diluted with 20 ml of water and brine and extracted with 3 x 80 ml ethyl acetate. The organic layers were washed four times with water and brine, dried over sodium sulfate, and evaporated to dryness. The crude product was purified by FLASH chromatography on a 30 mm x 5" silica gel column with hexane-ethyl acetate (3:2), and by HPLC on a silica gel column with hexane-ethyl acetate (1:1). It gave 90 mg (74%) of the title compound; m.p. 91-92 C, crystallization from methyl acetate-hexane.

[0115] Creams, gels and solutions containing ingredients within the proportions set forth in Examples A through C below, can be formulated by conventional means. Reference to Compound A, which is 1,25-dihydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol (not according to the invention), in these Examples is illustrative of any of the claimed compounds.

Example A

[0116]

5	Cream	
	Ingredients	% w/w
	Compound A	0.00001 - 0.10
10	Cetyl Alcohol	1.50
	Stearyl Alcohol	2.50
	Sorbitan Monostearate (Span 60)	2.00
	Mineral Oil	2.00
15	Glyceryl Monostearate and Polyoxy-ethylene Glycol Stearate Blend (Arlacel 165)	4.00
	Polysorbate 60 (Tween 60)	1.00
	Caprylic/Capric Triglyceride	5.00
20	Sorbitol Solution	4.00
	Edetate Disodium	0.10
	Butylated Hydroxyanisole (BHA)	0.02
	Sorbic Acid	0.20
25	Potassium Sorbate	0.1 - 0.2
	Water q.s. to	100.00

Example B

[0117]

35	Gel	
	Ingredients	% w/w
	Compound A	0.00001 - 0.10
	Butylated Hydroxyanisole (BHA)	0.02
40	Hydroxypropyl Cellulose	3.00
	Ethyl Alcohol, USP	45.00
	Water q.s. to	100.00

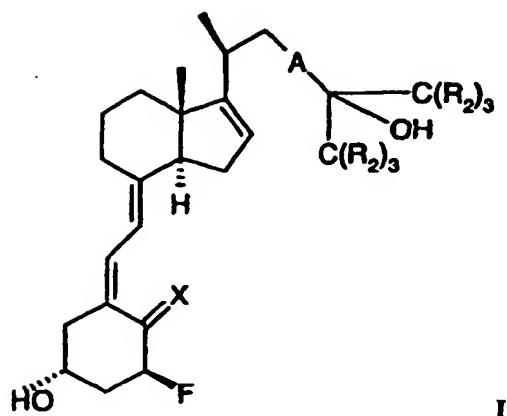
Example C

[0118]

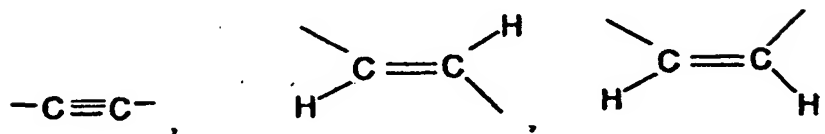
50	Topical Solution	
	Ingredients	% w/w
	Compound A	0.00001 - 0.10
	Propylene Glycol	10.00
55	Caprylic/Capric Triglyceride	30.00
	Butylated Hydroxyanisole (BHA)	0.02
	Ethyl Alcohol, Absolute q.s. to	100.00

Claims

1. A compound of the formula



wherein R_2 is hydrogen, C_{1-4} -alkyl or halogen, X is $=CH_2$,
and A is



or

$-CH_2-CH_2-$, provided that when A is $-CH_2-CH_2-$, R_2 is hydrogen or C_{1-4} -alkyl.

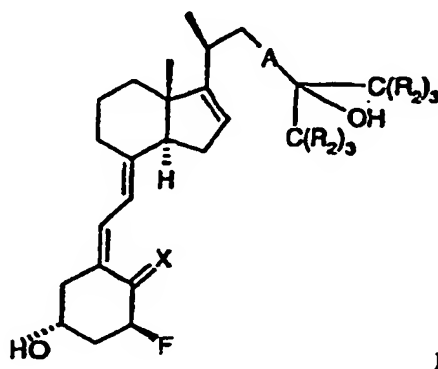
2. The compound of Claim 1, wherein R_2 is fluorine.
3. The compound of Claim 1, wherein R_2 is hydrogen.
4. The compound of Claim 1, wherein R_2 is C_{1-4} -alkyl.
5. The compound of Claim 1, wherein R_2 is methyl or ethyl.
6. The compounds:
 - 1 α -fluoro-25-hydroxy-16,23Z-diene-26,27-hexafluoro-20-epi-cholecalciferol,
 - 1 α -fluoro-25-hydroxy-16-ene-23-yne-26,27-hexafluoro-20-epi-cholecalciferol,
 - 1 α -fluoro-25-hydroxy-16,23E-diene-26,27-hexafluoro-20-epi-cholecalciferol,
 - 1 α -fluoro-25-hydroxy-16-ene-23-yne-26,27-bishomo-20-epi-cholecalciferol,
 - 1 α -fluoro-25-hydroxy-16,23E-diene-26,27-bishomo-20-epi-cholecalciferol.
7. A pharmaceutical composition comprising an effective amount of a compound of the formula I as in claim 1, and a pharmaceutically acceptable carrier.
8. A process for the manufacture of the compounds of formula I as in claim 1, which comprises deprotecting corresponding compounds containing silyl protected hydroxy groups.
9. The compounds of formula I as in claim 1, for use in inducing differentiation and inhibition of proliferation in skin

and cancer cell lines, particularly for treating hyperproliferative skin diseases, such as psoriasis; neoplastic diseases, such as leukemia; and sebaceous gland diseases, such as acne, for use in reversing the conditions associated with photodamage, particularly for the topical treatment of the skin damaged through sun exposure, the effects of wrinkling, elastosis and premature aging, and for use in the treatment of osteoporosis.

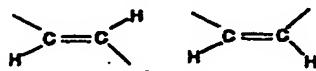
10. The use of the compounds of formula I as in claim 1 for the manufacture of pharmaceutical compositions
11. The use according to claim 10 for the manufacture of pharmaceutical compositions for inducing differentiation and inhibition of proliferation in skin and cancer cell lines, particularly for treating hyperproliferative skin diseases, such as psoriasis; neoplastic diseases, such as leukemia; and sebaceous gland diseases, such as acne.
12. The use according to claim 10 for the manufacture of pharmaceutical compositions for reversing the conditions associated with photodamage, particularly for the topical treatment of the skin damaged through sun exposure, the effects of wrinkling, elastosis and premature aging.
13. The use according to claim 10 for the manufacture of pharmaceutical compositions for the treatment of osteoporosis.

Patentansprüche

1. Verbindung der Formel



worin R_2 Wasserstoff, C_{1-4} -Alkyl oder Halogen darstellt, $X = CH_2$ darstellt, und $A = C \equiv C$,



oder $-CH_2-CH_2-$ darstellt, mit der Maßgabe, dass, wenn $A = -CH_2-CH_2-$ darstellt, R_2 Wasserstoff oder C_{1-4} -Alkyl darstellt.

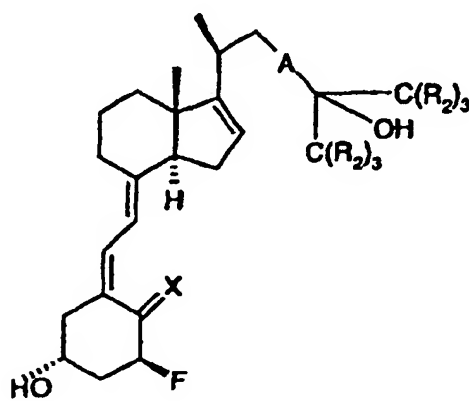
2. Verbindung nach Anspruch 1, worin R_2 Fluor darstellt.
3. Verbindung nach Anspruch 1, worin R_2 Wasserstoff darstellt.
4. Verbindung nach Anspruch 1, worin R_2 C_{1-4} -Alkyl darstellt.
5. Verbindung nach Anspruch 1, worin R_2 Methyl oder Ethyl darstellt.
6. Verbindungen:

1 α -Fluor-25-hydroxy-16,23Z-dien-26,27-hexafluor-20-epi-cholecalciferol,
1 α -Fluor-25-hydroxy-16-en-23-in-26,27-hexafluor-20-epi-cholecalciferol,
1 α -Fluor-25-hydroxy-16,23E-dien-26,27-hexafluor-20-epi-cholecalciferol,
1 α -Fluor-25-hydroxy-16-en-23-in-26,27-bishomo-20-epi-cholecalciferol,
1 α -Fluor-25-hydroxy-16,23E-dien-26,27-bishomo-20-epi-cholecalciferol.

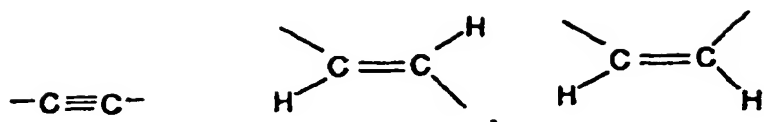
7. Pharmazeutische Zusammensetzung, umfassend eine wirksame Menge einer Verbindung der Formel I nach Anspruch 1 und einen pharmazeutisch verträglichen Träger.
8. Verfahren zur Herstellung von Verbindungen der Formel I nach Anspruch 1, das Schutzgruppenentfernung der entsprechenden Silyl-geschützte Hydroxygruppen enthaltenden Verbindungen umfasst.
9. Verbindungen der Formel I nach Anspruch 1 zur Verwendung bei der Induzierung von Differenziation und Inhibition von Proliferation in Haut- und Krebs-Zelllinien, insbesondere zum Behandeln von hyperproliferativen Hauterkrankungen, wie Psoriasis; neoplastischen Erkrankungen, wie Leukämie; und Talgdrüsenerkrankungen, wie Akne, zur Verwendung bei der Umkehrung der Zustände, die mit Lichtschädigung verbunden sind, insbesondere zur örtlichen Behandlung von durch Aussetzen der Sonne geschädigter Haut, den Wirkungen von Faltenbildung, Elastose und vorzeitigem Altern und zur Verwendung bei der Behandlung von Osteoporose.
10. Verwendung der Verbindungen der Formel I nach Anspruch 1 zur Herstellung von pharmazeutischen Zusammensetzungen.
11. Verwendung nach Anspruch 10 zur Herstellung von pharmazeutischen Zusammensetzungen zur Induzierung von Differenziation und Inhibition von Proliferation in Haut und Krebs-Zelllinien, insbesondere zum Behandeln von hyperproliferativen Hauterkrankungen, wie Psoriasis; neoplastischen Erkrankungen, wie Leukämie; und Talgdrüsenerkrankungen, wie Akne.
12. Verwendung nach Anspruch 10 zur Herstellung von pharmazeutischen Zusammensetzungen zur Umkehrung der Zustände, die mit Lichtschädigung verbunden sind, insbesondere zur örtlichen Behandlung von durch Aussetzen der Sonne geschädigter Haut, den Wirkungen von Faltenbildung, Elastose und vorzeitigem Altern.
13. Verwendung nach Anspruch 10 zur Herstellung von pharmazeutischen Zusammensetzungen zur Behandlung von Osteoporose.

Revendications

1. Composé de formule



dans lequel R₂ est un hydrogène, un alkyle en C₁₋₄ ou un halogène, X est =CH₂ et A est



or

10 ou $-\text{CH}_2-\text{CH}_2-$ à condition que lorsque A est $-\text{CH}_2-\text{CH}_2-$, R_2 est un hydrogène ou un alkyle en C_{1-4} .

2. Composé selon la revendication 1, dans lequel R_2 est le fluor.

3. Composé selon la revendication 1, dans lequel R_2 est l'hydrogène.

4. Composé selon la revendication 1, dans lequel R_2 est un alkyle en C_{1-4} .

5. Composé selon la revendication 1, dans lequel R_2 est un groupement méthyle ou éthyle.

6. Composés :

1 α -fluoro-25-hydroxy-16,23Z-diène-26,27-hexafluoro-20-épi-cholécalciférol
 1 α -fluoro-25-hydroxy-16-ène-23-yne-26,27-hexafluoro-20-épi-cholécalciférol
 1 α -fluoro-25-hydroxy-16,23E-diène-26,27-hexafluoro-20-épi-cholécalciférol
 1 α -fluoro-25-hydroxy-16-ène-23-yne-26,27-bishomo-20-épi-cholécalciférol
 1 α -fluoro-25-hydroxy-16,23E-diène-26,27-bishomo-20-épi-cholécalciférol

7. Composition pharmaceutique comprenant une quantité effective d'un composé de la formule I selon la revendication 1, et un support pharmaceutiquement acceptable.

8. Procédé pour la fabrication des composés de formule I selon la revendication 1, qui comprend des composés correspondants de déprotection contenant des groupes hydroxy protégés par un silyle.

9. Composés de formule I selon la revendication 1, à utiliser pour induire la différenciation et l'inhibition de la prolifération dans la peau et les lignes cellulaires cancéreuses, en particulier pour traiter les maladies de la peau hyperprolifératives, comme le psoriasis ; les maladies néoplasiques, comme la leucémie ; et les maladies des glandes sébacées, comme l'acné, à utiliser dans l'inversion des conditions associées au photodommage, particulièrement pour le traitement topique de la peau endommagée par l'exposition au soleil, les effets du plissement, la dégénérescence élastique et le vieillissement prématuré, et à utiliser dans le traitement de l'ostéoporose.

10. Utilisation des composés de formule I selon la revendication 1, pour la fabrication de compositions pharmaceutiques.

11. Utilisation selon la revendication 10 pour la fabrication de compositions pharmaceutiques afin d'induire une différenciation et une inhibition de la prolifération dans la peau et les lignes cellulaires cancéreuses, particulièrement pour le traitement des maladies de la peau hyperprolifératives, comme le psoriasis ; les maladies néoplastiques comme la leucémie, et les maladies des glandes sébacées, comme l'acné.

12. Utilisation selon la revendication 10, pour la fabrication de compositions pharmaceutiques pour inverser les conditions associées au photodommage, particulièrement pour le traitement topique de la peau endommagée par l'exposition au soleil, les effets du plissement, la dégénérescence élastique et le vieillissement prématuré.

13. Utilisation selon la revendication 10, pour la fabrication de compositions pharmaceutiques pour le traitement de l'ostéoporose.